

## PEPTIDES WHICH TARGET TUMOR AND ENDOTHELIAL CELLS, COMPOSITIONS AND USES THEREOF

This application claims the benefit under 35 U.S.C. § 119(e) from United  
5 States Provisional Application Serial No. 60/429,174, filed November 25, 2002 and  
United States Provisional Application Serial No. 60/475,539, filed June 2, 2003 which  
are herein incorporated by reference in their entirety.

### 1. Field of the Invention

10 The present invention relates generally to peptide analogs of Ac-PHSCN-NH<sub>2</sub>  
which target tumor and endothelial cells and have anti-tumor, anti-angiogenic and  
anti-metastatic activity, methods of making these peptides, compositions thereof and  
methods of using these peptides and pharmaceutical compositions thereof to treat,  
prevent and detect diseases characterized by tumor growth, metastasis and  
15 angiogenesis. The peptide analogs may serve, *inter alia*, as carriers of radioactivity,  
PET-active compounds, toxins, fluorescent molecules and PEG molecules.

### 2. Background of the Invention

Integrins are heterodimeric, transmembrane proteins that are involved in cell  
20 adhesion, motility and survival (Geiger *et al.*, *Nat. Rev. Mol. Cell Biol* **2001**,  
2(11):793-805). Integrin ligands comprise the extracellular matrix (ECM) and  
basement membranes and include collagen, laminin, fibronectin, vitronectin, and  
fibrinogen (Bokel, *Dev. Cell* **2002**, 3(3):311-21; Stupack *et al.*, *J Cell Sci* **2002**, 115:  
3729-38; Bornstein *et al.*, *Curr Opin Cell Biol* **2002**, 14(5): 608-16). Integrin  
25 expression and importantly, integrin activation state is altered in tumor tissue (Liu *et al.*,  
*Semin Oncol* **2002**, 29 (3 Suppl 11): 96-103; Hood *et al.*, *Nat. Rev. Cancer* **2002**  
2(2): 91-100; Felding-Habermann, *Clin Exp Metastasis* **2003**, 20(3): 203-13).  
Accordingly, several integrins, including  $\alpha_v\beta_3$ ,  $\alpha_v\beta_5$  and  $\alpha_5\beta_1$ , have been recognized as  
validated therapeutic targets for the treatment and prevention of cancer (Kumar, *Curr*  
30 *Drug Targets* **2003**, 4(2): 123-31; Kerr *et al.*, *Expert Opin Investig Drugs* **2002**,  
11(12): 1765-74; Rust *et al.*, *J Biomed Biotechnol.* **2002**, 2(3): 124-130; Damiano,  
*Curr Cancer Drug Targets* **2002**, 2(1): 37-43; Tucker, *Curr Opin Pharmacol* **2002**,  
2(4): 394-402) and receptor-targeted imaging methods for oncology (Herschman,

*Science* **2003**, 302(5645): 605-8); Aboagye *et al.*, *Invest New Drugs* **2003**, 21(2): 169-81; Van De Wiele *et al.*, *Eur J Nucl Med Mol Imaging* **2002** 29(5): 699-709; Glaser *et al.*, *Int J Oncol* **2003**, 22(2):253-67).

The integrin  $\alpha_5\beta_1$  is normally not expressed in quiescent endothelial cells, but  
5 is upregulated during angiogenesis (Kim *et al.*, *Am J Pathol* **2000**, 156(4):1345-62).  
 $\alpha_5\beta_1$  is a receptor for fibronectin, an abundant plasma protein that may also be  
associated with the extracellular matrix (Labat-Robert, *Semin Cancer Biol* **2002**,  
12(3): 187-95). Fibronectin interacts with the  $\alpha_5\beta_1$  integrin through several epitopes  
with the major adhesive interaction mediated by the RGD sequence within the 10<sup>th</sup>  
10 type III repeat. The major adhesive interaction mediates cell signaling events and can  
be potentiated by a second epitope located in the ninth Type III repeat called the  
synergy region (*i.e.*, PHSRN) (Akiyama *et al.*, *Cancer Metastasis Rev* **1995**, 14(3):  
173-89). Antagonists of  $\alpha_5\beta_1$  were able to inhibit tumor angiogenesis and cause  
tumor regression which demonstrates the therapeutic potential of targeting the integrin  
15  $\alpha_5\beta_1$  (Kim *et al.*, *supra*). The integrin  $\alpha_5\beta_1$  also appears to be important in the  
survival and metastasis of tumor cells (O'Brien *et al.*, *Exp Cell Res* **1996**, 224(1): 208-  
13; Ruoslahti, *Invasion Metastasis* 1994-95;14(1-6):87-9724-26); Kemperman *et al.*,  
*Invasion Metastasis* 1994-95,14(1-6): 98-108; Tani *et al.*, *Br J Cancer* **2003**, 88(2):  
327-33).

20 The integrin  $\alpha_v\beta_3$  is also a therapeutic target for the inhibition of tumor  
angiogenesis since inhibitors of  $\alpha_v\beta_3$  (*e.g.*, monoclonal antibodies, cyclic RGD  
peptides and small non-peptidic organic compounds) are efficacious in multiple pre-  
clinical models of cancer progression (Kumar, *Curr Drug Targets* **2003**, 4(2): 123-31;  
Varner *et al.*, *Important Adv Oncol* **1996**, 69-87; Brooks, *J Clin Invest* **1995**, 96(4):  
25 1815-22). Although  $\alpha_v\beta_3$  is not normally expressed on epithelial cells, it is up-  
regulated on tumor cells, leading to tumor cell adhesion, migration and invasion  
(Metzner *et al.*, *J Invest Dermatol* **1996**, 107(4): 597-602). The  $\alpha_v\beta_3$  integrin has  
been implicated in melanoma progression and metastasis (Nip *et al.*, *Cancer*  
*Metastasis Rev* **1995**, 14(3): 241-52) and  $\alpha_v\beta_3$  expression has been documented in a  
30 variety of tumor cell types including breast, prostate, pancreas, kidney, and glioma  
(Felding-Habermann *et al.*, *Proc Natl Acad Sci USA* **2001**, 98(4): 1853-8; Platten *et al.*,  
*Biochem Biophys Res Commun* **2000**, 268(2): 607-11; Lohr *et al.*, *Pancreas* **1996**,  
12(3): 248-59; Rabb *et al.*, *Am J Nephrol* **1996**, 16(5): 402-8; Cooper *et al.*, *Neoplasia*

2002, 4(3): 191-4). Further, the expression of  $\alpha_v\beta_3$  has also been associated with metastasis to bone (Pecheur *et al.*, *FASEB J* 2002, 16(10): 1266-8.).

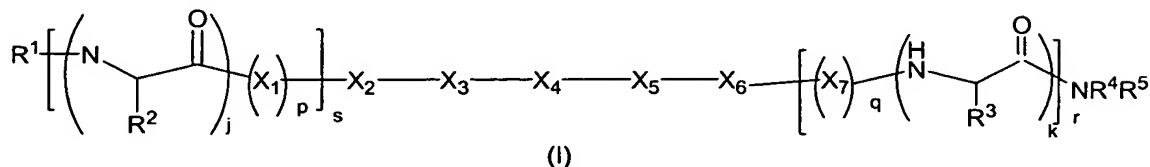
Ac-PHSCN-NH<sub>2</sub> is derived from the synergy sequence of fibronectin and has been shown to target activated  $\alpha_5\beta_1$  and  $\alpha_V\beta_3$  integrins on cell surfaces (Livant, United States Patent No. 6,001,965; Livant, United States Patent No. 6,472,369; Livant *et al.*, *Cancer Res* 2000, 60(2): 309-20). Further, Ac-PHSCN-NH<sub>2</sub> completely inhibits DU145 invasion and metastasis of MatLyLu cells in a rat model (Livant *et al.*, *supra*) and in combination with 5-FU infusion improved survival in a CT26 colon cancer model (Stoeltzing *et al.*, *Int J Cancer* 2003, 104(4): 496-503). Accordingly, Ac-PHSCN-NH<sub>2</sub> targets tumors and the blood vessels which nourish tumor cells as shown by visualization of a Ac-PHSCN-NH<sub>2</sub> derivative.

Thus, what is needed are novel peptide analogs of Ac-PHSCN-NH<sub>2</sub> to fully explore the potential of Ac-PHSCN-NH<sub>2</sub> derivatives in targeting tumors and the vasculature which nourishes tumor cells. Ideally, the peptide analogs will serve, *inter alia*, as carriers of radioactivity for imaging and radiotherapy, PET-active compounds for PET-imaging, toxins for targeted delivery of cellular toxins, fluorescent molecules for visualization and PEG molecules for improvement of pharmacokinetic parameters.

### 3. Summary of the Invention

The present invention satisfies these and other needs by providing peptide analogs of Ac-PHSCN-NH<sub>2</sub> which target tumor and endothelial cells and have anti-tumor, anti-angiogenic and anti-metastatic activity, methods of making these peptides, compositions thereof and methods of using these peptides and pharmaceutical compositions thereof to treat, prevent and detect diseases characterized by tumor growth, metastasis and angiogenesis. The peptide analogs may serve, *inter alia*, as carriers of radioactivity, PET-active compounds, toxins, fluorescent molecules and PEG molecules.

In one aspect the present invention provides a compound of Formula (I):



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or a pharmaceutically acceptable salt, solvate, hydrate or N-oxide thereof  
wherein:

j and k are independently 0 or 1;

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p and q are independently an integer including and between 0 and 100;

r and s are independently 0 or 1;

10  $R^1$  is acyl, substituted acyl, acyl chelate, alkyl, substituted alkyl, cycloalkyl, substituted cycloalkyl, imino or substituted imino;

$R^2$  is  $C_1$ - $C_6$  alkyl with at least one hydrogen atom replaced by a substituent selected from the group consisting of  $-NR^6R^7$ ,  $-OR^8$ ,  $-CO_2R^9$ ,  $-S(O)_zR^{10}$ , -

15  $P(OR^{11})OR^{12}$ , aryl and substituted aryl;

$R^6$ ,  $R^7$ ,  $R^8$ ,  $R^9$ ,  $R^{10}$ ,  $R^{11}$  and  $R^{12}$  are independently selected from the group consisting of hydrogen, acyl, substituted acyl, acyl chelate, alkyl, substituted alkyl, cycloalkyl, substituted cycloalkyl, imino and substituted imino;

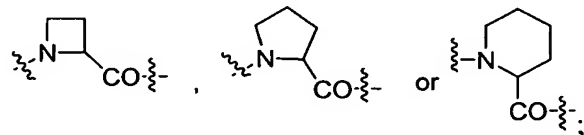
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$X_1$  is  $-NH(C=C)_gCO-$ ,  $-NH(CH_2)_hCO-$  or  $-NHCH(CH_3)CO-$ ;

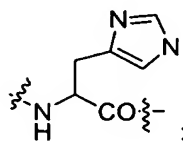
g and h are independently 1, 2, 3, 4, 5 or 6

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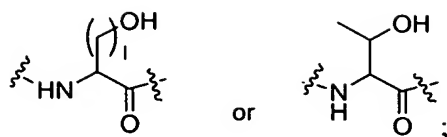
$X_2$  is



$X_3$  is



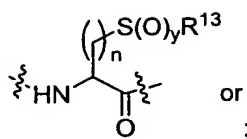
X<sub>4</sub> is



l is an integer from 1 to 4;

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X<sub>5</sub> is



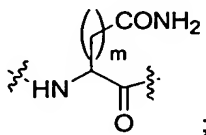
R<sup>13</sup> is hydrogen, alkyl, substituted alkyl, acyl, substituted acyl, arylalkyl,  
10 substituted arylalkyl, aryl, substituted aryl or -S(O)<sub>x</sub>R<sup>14</sup>;

n is an integer from 1 to 5;

R<sup>14</sup> is alkyl, substituted alkyl, acyl, substituted acyl, arylalkyl, substituted  
15 arylalkyl, aryl or substituted aryl;

y and x are independently 0, 1 or 2;

X<sub>6</sub> is



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m is an integer from 1, 2, 3 or 4;

X<sub>7</sub> is -NH(C=C)<sub>d</sub>CO-, -NH(CH<sub>2</sub>)<sub>e</sub>CO- or -NHCH(CH<sub>3</sub>)CO-;

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d and e are independently 1, 2, 3, 4, 5 or 6;

R<sup>3</sup> is C<sub>1</sub>-C<sub>6</sub> alkyl with at least one hydrogen atom replaced by a substituent selected from the group consisting of -NR<sup>15</sup>R<sup>16</sup>, -OR<sup>17</sup>, -CO<sub>2</sub>R<sup>18</sup>, -S(O)<sub>n</sub>R<sup>19</sup>, -P(OR<sup>20</sup>)OR<sup>21</sup>, aryl and substituted aryl;

5 R<sup>4</sup> and R<sup>5</sup> are independently hydrogen, alkyl or substituted alkyl; and

R<sup>15</sup>, R<sup>16</sup>, R<sup>17</sup>, R<sup>18</sup>, R<sup>19</sup>, R<sup>20</sup> and R<sup>21</sup> are independently selected from the group consisting of hydrogen, acyl, substituted acyl, acyl chelate, alkyl, substituted alkyl, cycloalkyl, substituted cycloalkyl, imino and substituted imino;

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with the proviso that R<sup>1</sup> is not acetyl when R<sup>4</sup> and R<sup>5</sup> are hydrogen and r and s are 0.

In a second aspect, the present invention provides pharmaceutical compositions of compounds of the invention. The pharmaceutical compositions generally comprise one or more compounds of the invention or pharmaceutically acceptable salts, hydrates or solvates thereof and a pharmaceutically acceptable diluent, carrier, excipient and adjuvant. The choice of diluent, carrier, excipient and adjuvant will depend upon, among other factors, the desired mode of administration.

15 In a third aspect, the present invention provides methods for treating or preventing diseases or disorders such as cancer. The methods generally involve administering to a patient in need of such treatment or prevention a therapeutically effective amount of a compound of the invention and/or a pharmaceutical composition thereof.

20 In a fourth aspect, the present invention provides methods for detecting diseases or disorders such as cancer. The methods generally involve administering to a patient in need of such treatment or prevention a diagnostically effective amount of a compound of the invention and/or a pharmaceutical composition thereof.

#### 4. Detailed Description Of The Invention

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##### 4.1 Definitions

“Compounds of the invention” refers to compounds encompassed by structural formulae (I), (II), (III), (IV) and (V) disclosed herein and includes any specific compounds within that generic formula whose structure is disclosed herein. The compounds of the invention may be identified either by their chemical structure

and/or chemical name. When the chemical structure and chemical name conflict, the chemical structure is determinative of the identity of the compound. The compounds of the invention may contain one or more chiral centers and/or double bonds and therefore, may exist as stereoisomers, such as double-bond isomers (*i.e.*, geometric isomers), enantiomers or diastereomers. Accordingly, the chemical structures depicted herein encompass all possible enantiomers and stereoisomers of the illustrated compounds including the stereoisomerically pure form (*e.g.*, geometrically pure, enantiomerically pure or diastereomerically pure) and enantiomeric and stereoisomeric mixtures. The compounds of the invention may also exist in several tautomeric forms. Accordingly, the chemical structures depicted herein encompass all possible tautomeric forms of the illustrated compounds. The compounds of the invention also include isotopically labeled compounds where one or more atoms have an atomic mass different from the atomic mass conventionally found in nature. Examples of isotopes that may be incorporated into the compounds of the invention include, but are not limited to,  $^2\text{H}$ ,  $^3\text{H}$ ,  $^{13}\text{C}$ ,  $^{14}\text{C}$ ,  $^{15}\text{N}$ ,  $^{17}\text{O}$ ,  $^{18}\text{O}$ , *etc.* Compounds of the invention may exist in unsolvated forms as well as solvated forms, including hydrated forms and as N-oxides. In general, the hydrated, solvated and N-oxide forms are within the scope of the present invention. Certain compounds of the present invention may exist in multiple crystalline or amorphous forms. In general, all physical forms are equivalent for the uses contemplated by the present invention and are intended to be within the scope of the present invention. Further, it should be understood, when partial structures of the compounds of the invention are illustrated, that brackets indicate the point of attachment of the partial structure to the rest of the molecule.

“Alkyl” by itself or as part of another substituent refers to a saturated or unsaturated, branched, straight-chain or cyclic monovalent hydrocarbon radical derived by the removal of one hydrogen atom from a single carbon atom of a parent alkane, alkene or alkyne. Typical alkyl groups include, but are not limited to, methyl; ethyls such as ethanyl, ethenyl, ethynyl; propyls such as propan-1-yl, propan-2-yl, cyclopropan-1-yl, prop-1-en-1-yl, prop-1-en-2-yl, prop-2-en-1-yl (allyl), cycloprop-1-en-1-yl; cycloprop-2-en-1-yl, prop-1-yn-1-yl, prop-2-yn-1-yl, *etc.*; butyls such as butan-1-yl, butan-2-yl, 2-methyl-propan-1-yl, 2-methyl-propan-2-yl, cyclobutan-1-yl, but-1-en-1-yl, but-1-en-2-yl, 2-methyl-prop-1-en-1-yl, but-2-en-1-yl, but-2-en-2-yl, buta-1,3-dien-1-yl, buta-1,3-dien-2-yl, cyclobut-1-en-1-yl,

cyclobut-1-en-3-yl, cyclobuta-1,3-dien-1-yl, but-1-yn-1-yl, but-1-yn-3-yl, but-3-yn-1-yl, *etc.*; and the like.

The term “alkyl” is specifically intended to include groups having any degree or level of saturation, *i.e.*, groups having exclusively single carbon-carbon bonds, groups having one or more double carbon-carbon bonds, groups having one or more triple carbon-carbon bonds and groups having mixtures of single, double and triple carbon-carbon bonds. Where a specific level of saturation is intended, the expressions “alkanyl,” “alkenyl,” and “alkynyl” are used. Preferably, an alkyl group comprises from 1 to 20 carbon atoms, more preferably, from 1 to 10 carbon atoms. (C<sub>1</sub>-C<sub>6</sub>) alkyl, for example, refers to an alkyl group containing from 1 to 6 carbon atoms.

“Alkanyl” by itself or as part of another substituent refers to a saturated branched, straight-chain or cyclic alkyl radical derived by the removal of one hydrogen atom from a single carbon atom of a parent alkane. Typical alkanyl groups include, but are not limited to, methanyl; ethanyl; propanyls such as propan-1-yl, propan-2-yl (isopropyl), cyclopropan-1-yl, *etc.*; butanyls such as butan-1-yl, butan-2-yl (*sec*-butyl), 2-methyl-propan-1-yl (isobutyl), 2-methyl-propan-2-yl (*t*-butyl), cyclobutan-1-yl, *etc.*; and the like.

“Alkenyl” by itself or as part of another substituent refers to an unsaturated branched, straight-chain or cyclic alkyl radical having at least one carbon-carbon double bond derived by the removal of one hydrogen atom from a single carbon atom of a parent alkene. The group may be in either the *cis* or *trans* conformation about the double bond(s). Typical alkenyl groups include, but are not limited to, ethenyl; propenyls such as prop-1-en-1-yl, prop-1-en-2-yl, prop-2-en-1-yl (allyl), prop-2-en-2-yl, cycloprop-1-en-1-yl; cycloprop-2-en-1-yl; butenyls such as but-1-en-1-yl, but-1-en-2-yl, 2-methyl-prop-1-en-1-yl, but-2-en-1-yl, but-2-en-1-yl, but-2-en-2-yl, buta-1,3-dien-1-yl, buta-1,3-dien-2-yl, cyclobut-1-en-1-yl, cyclobut-1-en-3-yl, cyclobuta-1,3-dien-1-yl, *etc.*; and the like.

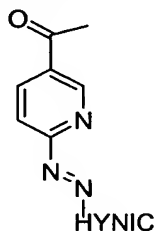
“Alkynyl” by itself or as part of another substituent refers to an unsaturated branched, straight-chain or cyclic alkyl radical having at least one carbon-carbon triple bond derived by the removal of one hydrogen atom from a single carbon atom of a parent alkyne. Typical alkynyl groups include, but are not limited to, ethynyl;



propynyls such as prop-1-yn-1-yl, prop-2-yn-1-yl, *etc.*; butynyls such as but-1-yn-1-yl, but-1-yn-3-yl, but-3-yn-1-yl, *etc.*; and the like.

5       “Acyl” by itself or as part of another substituent refers to a radical  $-C(O)R^{30}$ , where  $R^{30}$  is hydrogen, alkyl, cycloalkyl, cycloheteroalkyl, aryl, arylalkyl, heteroalkyl, heteroaryl, heteroarylalkyl as defined herein. Representative examples include, but are not limited to, formyl, acetyl, cyclohexylcarbonyl, cyclohexylmethylcarbonyl, benzoyl, benzylcarbonyl and the like.

10       “Acyl Chelate” by itself or as part of another substituent refers to a radical  $-C(O)R^{31}$ , where  $R^{31}$  is alkyl, cycloalkyl, aryl as defined herein substituted with a chelating group which binds an appropriate metal. Representative examples include  $-C(O)CH_2CH_2-R^{32}$  where  $R^{32}$  is a chelating group such as, for example, DOTA, TETA, a polyamino carboxylate (*e.g.*, NODAGA, EDTA, tricine,  $-C(O)CH_2-DTPA$ , *etc.*)



15       and where the chelating group is bound to metals such as positron emitting labels (*e.g.*,  $^{18}F$ ,  $^{45}Ti$ ,  $^{44}Sc$ ,  $^{55}Co$ ,  $^{61}Cu$ ,  $^{64}Cu$ ,  $^{66}Ga$ ,  $^{68}Ga$ ,  $^{75}Br$ ,  $^{76}Br$ ,  $^{86}Y$ ,  $^{110}In$ ,  $^{124}I$ ,  $^{89}Zr$ ,  $^{99}Tc$ ), radionuclides (*e.g.*,  $^{137}Cs$ ,  $^{60}Co$ ,  $^{131}I$ ,  $^{123}I$ ,  $^{192}Ir$ ,  $^{90}Y$ ,  $^{67}Ga$ ,  $^{99}Tc$ ,  $^{123}I$ ,  $^{125}I$ ,  $^{131}I$ ,  $^{111}In$ ,  $^{97}Ru$ ,  $^{67}Cu$ ,  $^{68}Ga$ ,  $^{72}As$ ,  $^{89}Zr$ ,  $^{90}Y$ ,  $^{201}Tl$ , *etc.*) or lanthanide metals.

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      “Alkoxy” by itself or as part of another substituent refers to a radical  $-OR^{31}$  where  $R^{33}$  represents an alkyl or cycloalkyl group as defined herein. Representative examples include, but are not limited to, methoxy, ethoxy, propoxy, butoxy, cyclohexyloxy and the like.

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      “Aryl” by itself or as part of another substituent refers to a monovalent aromatic hydrocarbon radical derived by the removal of one hydrogen atom from a single carbon atom of a parent aromatic ring system. Typical aryl groups include, but are not limited to, groups derived from aceanthrylene, acenaphthylene, acphenanthrylene, anthracene, azulene, benzene, chrysene, coronene, fluoranthene,

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fluorene, hexacene, hexaphene, hexalene, *as*-indacene, *s*-indacene, indane, indene, naphthalene, octacene, octaphene, octalene, ovalene, penta-2,4-diene, pentacene, pentalene, pentaphene, perylene, phenalene, phenanthrene, picene, pleiadene, pyrene, pyranthrene, rubicene, triphenylene, trinaphthalene and the like. Preferably, an aryl group comprises from 6 to 20 carbon atoms, more preferably from 6 to 12 carbon atoms.

“Arylalkyl” by itself or as part of another substituent refers to an acyclic alkyl radical in which one of the hydrogen atoms bonded to a carbon atom, typically a terminal or  $sp^3$  carbon atom, is replaced with an aryl group. Typical arylalkyl groups include, but are not limited to, benzyl, 2-phenylethan-1-yl, 2-phenylethen-1-yl, naphthylmethyl, 2-naphthylethan-1-yl, 2-naphthylethen-1-yl, naphthobenzyl, 2-naphthophenylethan-1-yl and the like. Where specific alkyl moieties are intended, the nomenclature arylalkanyl, arylalkenyl and/or arylalkynyl is used. Preferably, an arylalkyl group is ( $C_6$ - $C_{30}$ ) arylalkyl, *e.g.*, the alkanyl, alkenyl or alkynyl moiety of the arylalkyl group is ( $C_1$ - $C_{10}$ ) and the aryl moiety is ( $C_6$ - $C_{20}$ ), more preferably, an arylalkyl group is ( $C_6$ - $C_{20}$ ) arylalkyl, *e.g.*, the alkanyl, alkenyl or alkynyl moiety of the arylalkyl group is ( $C_1$ - $C_8$ ) and the aryl moiety is ( $C_6$ - $C_{12}$ ).

“Cycloalkyl” by itself or as part of another substituent refers to a saturated or unsaturated cyclic alkyl radical. Where a specific level of saturation is intended, the nomenclature “cycloalkanyl” or “cycloalkenyl” is used. Typical cycloalkyl groups include, but are not limited to, groups derived from cyclopropane, cyclobutane, cyclopentane, cyclohexane and the like. Preferably, the cycloalkyl group is ( $C_3$ - $C_{10}$ ) cycloalkyl, more preferably ( $C_3$ - $C_7$ ) cycloalkyl.

“Cycloheteroalkyl” by itself or as part of another substituent refers to a saturated or unsaturated cyclic alkyl radical in which one or more carbon atoms (and any associated hydrogen atoms) are independently replaced with the same or different heteroatom. Typical heteroatoms to replace the carbon atom(s) include, but are not limited to, N, P, O, S, Si, *etc.* Where a specific level of saturation is intended, the nomenclature “cycloheteroalkanyl” or “cycloheteroalkenyl” is used. Typical cycloheteroalkyl groups include, but are not limited to, groups derived from epoxides,

azirines, thiiranes, imidazolidine, morpholine, piperazine, piperidine, pyrazolidine, pyrrolidine, quinuclidine, and the like.

5       “Diagnostically effective amount” means the amount of a compound that, when administered to a patient for detection of a disease, is sufficient to detect the disease. The “diagnostically effective amount” will vary depending on the compound, the disease and its severity and the age, weight, *etc.*, of the patient to be treated.

10       “Heteroalkyl, Heteroalkanyl, Heteroalkenyl and Heteroalkynyl” by themselves or as part of another substituent refer to alkyl, alkanyl, alkenyl and alkynyl groups, respectively, in which one or more of the carbon atoms (and any associated hydrogen atoms) are independently replaced with the same or different heteroatomic groups. Typical heteroatomic groups which can be included in these groups include, but are  
15       not limited to, -O-, -S-, -O-O-, -S-S-, -O-S-, -NR<sup>34</sup>R<sup>35</sup>, -N=N-, -N=N-, -N=N-NR<sup>36</sup>R<sup>37</sup>, -PR<sup>38</sup>-, -P(O)<sub>2</sub>-, -POR<sup>39</sup>-, -O-P(O)<sub>2</sub>-, -SO-, -SO<sub>2</sub>-, -SnR<sup>40</sup>R<sup>41</sup>- and the like, where R<sup>34</sup>, R<sup>35</sup>, R<sup>36</sup>, R<sup>37</sup>, R<sup>38</sup>, R<sup>39</sup>, R<sup>40</sup> and R<sup>41</sup> are independently hydrogen, alkyl, substituted alkyl, aryl, substituted aryl, arylalkyl, substituted arylalkyl, cycloalkyl, substituted cycloalkyl, cycloheteroalkyl, substituted cycloheteroalkyl, heteroalkyl,  
20       substituted heteroalkyl, heteroaryl, substituted heteroaryl, heteroarylalkyl or substituted heteroarylalkyl.

      “Heteroaryl” by itself or as part of another substituent refers to a monovalent heteroaromatic radical derived by the removal of one hydrogen atom from a single  
25       atom of a parent heteroaromatic ring system. Typical heteroaryl groups include, but are not limited to, groups derived from acridine, arsindole, carbazole,  $\beta$ -carboline, chromane, chromene, cinnoline, furan, imidazole, indazole, indole, indoline, indolizine, isobenzofuran, isochromene, isoindole, isoindoline, isoquinoline, isothiazole, isoxazole, naphthyridine, oxadiazole, oxazole, perimidine,  
30       phenanthridine, phenanthroline, phenazine, phthalazine, pteridine, purine, pyran, pyrazine, pyrazole, pyridazine, pyridine, pyrimidine, pyrrole, pyrrolizine, quinazoline, quinoline, quinolizine, quinoxaline, tetrazole, thiadiazole, thiazole, thiophene, triazole, xanthene, and the like. Preferably, the heteroaryl group is from 5-20

membered heteroaryl, more preferably from 5-10 membered heteroaryl. Preferred heteroaryl groups are those derived from thiophene, pyrrole, benzothiophene, benzofuran, indole, pyridine, quinoline, imidazole, oxazole and pyrazine

5           “Heteroarylalkyl” by itself or as part of another substituent refers to an acyclic alkyl radical in which one of the hydrogen atoms bonded to a carbon atom, typically a terminal or  $sp^3$  carbon atom, is replaced with a heteroaryl group. Where specific alkyl moieties are intended, the nomenclature heteroarylalkanyl, heteroarylalkenyl and/or heteroarylalkynyl is used. In preferred embodiments, the heteroarylalkyl group is a  
10 6-30 membered heteroarylalkyl, *e.g.*, the alkanyl, alkenyl or alkynyl moiety of the heteroarylalkyl is 1-10 membered and the heteroaryl moiety is a 5-20-membered heteroaryl, more preferably, 6-20 membered heteroarylalkyl, *e.g.*, the alkanyl, alkenyl or alkynyl moiety of the heteroarylalkyl is 1-8 membered and the heteroaryl moiety is a 5-12-membered heteroaryl.

15

          “Imino” by itself or as part of another substituent refers to a radical  $-C=NR^{42}$ , where  $R^{42}$  is hydrogen, alkyl, cycloalkyl, cycloheteroalkyl, aryl, arylalkyl, heteroalkyl, heteroaryl, heteroarylalkyl as defined herein. .

20           “Parent Aromatic Ring System” refers to an unsaturated cyclic or polycyclic ring system having a conjugated  $\pi$  electron system. Specifically included within the definition of “parent aromatic ring system” are fused ring systems in which one or more of the rings are aromatic and one or more of the rings are saturated or unsaturated, such as, for example, fluorene, indane, indene, phenalene, *etc.* Typical  
25 parent aromatic ring systems include, but are not limited to, aceanthrylene, acenaphthylene, acephenanthrylene, anthracene, azulene, benzene, chrysene, coronene, fluoranthene, fluorene, hexacene, hexaphene, hexalene, *as*-indacene, *s*-indacene, indane, indene, naphthalene, octacene, octaphene, octalene, ovalene, penta-2,4-diene, pentacene, pentalene, pentaphene, perylene, phenalene,  
30 phenanthrene, picene, pleiadene, pyrene, pyranthrene, rubicene, triphenylene, trinaphthalene and the like.

“Parent Heteroaromatic Ring System” refers to a parent aromatic ring system in which one or more carbon atoms (and any associated hydrogen atoms) are independently replaced with the same or different heteroatom. Typical heteroatoms to replace the carbon atoms include, but are not limited to, N, P, O, S, Si, *etc.*

5 Specifically included within the definition of “parent heteroaromatic ring systems” are fused ring systems in which one or more of the rings are aromatic and one or more of the rings are saturated or unsaturated, such as, for example, arsindole, benzodioxan, benzofuran, chromane, chromene, indole, indoline, xanthene, *etc.* Typical parent heteroaromatic ring systems include, but are not limited to, arsindole, carbazole,

10  $\beta$ -carboline, chromane, chromene, cinnoline, furan, imidazole, indazole, indole, indoline, indolizine, isobenzofuran, isochromene, isoindole, isoindoline, isoquinoline, isothiazole, isoxazole, naphthyridine, oxadiazole, oxazole, perimidine, phenanthridine, phenanthroline, phenazine, phthalazine, pteridine, purine, pyran, pyrazine, pyrazole, pyridazine, pyridine, pyrimidine, pyrrole, pyrrolizine, quinazoline,

15 quinoline, quinolizine, quinoxaline, tetrazole, thiadiazole, thiazole, thiophene, triazole, xanthene, and the like.

“Patient” includes humans. The terms “human” and “patient” are used interchangeably herein.

20

“Pharmaceutically acceptable salt” refers to a salt of a compound of the invention, which is pharmaceutically acceptable and possesses the desired pharmacological activity of the parent compound. Such salts include: (1) acid addition salts, formed with inorganic acids such as hydrochloric acid, hydrobromic

25 acid, sulfuric acid, nitric acid, phosphoric acid, and the like; or formed with organic acids such as acetic acid, propionic acid, hexanoic acid, cyclopentanepropionic acid, glycolic acid, pyruvic acid, lactic acid, malonic acid, succinic acid, malic acid, maleic acid, fumaric acid, tartaric acid, citric acid, benzoic acid, 3-(4-hydroxybenzoyl) benzoic acid, cinnamic acid, mandelic acid, methanesulfonic acid, ethanesulfonic

30 acid, 1,2-ethane-disulfonic acid, 2-hydroxyethanesulfonic acid, benzenesulfonic acid, 4-chlorobenzenesulfonic acid, 2-naphthalenesulfonic acid, 4-toluenesulfonic acid, camphorsulfonic acid, 4-methylbicyclo[2.2.2]-oct-2-ene-1-carboxylic acid, glucoheptonic acid, 3-phenylpropionic acid, trimethylacetic acid, tertiary butylacetic

acid, lauryl sulfuric acid, gluconic acid, glutamic acid, hydroxynaphthoic acid, salicylic acid, stearic acid, muconic acid, and the like; or (2) salts formed when an acidic proton present in the parent compound is replaced by a metal ion, *e.g.*, an alkali metal ion, an alkaline earth ion, or an aluminum ion; or coordinates with an organic  
5 base such as ethanolamine, diethanolamine, triethanolamine, N-methylglucamine and the like.

“Pharmaceutically acceptable vehicle” refers to a diluent, adjuvant, excipient or carrier with which a compound of the invention is administered.

10

“Preventing” or “prevention” refers to a reduction in risk of acquiring a disease or disorder (*i.e.*, causing at least one of the clinical symptoms of the disease not to develop in a patient that may be exposed to or predisposed to the disease but does not yet experience or display symptoms of the disease).

15

“Substituted” refers to a group in which one or more hydrogen atoms are independently replaced with the same or different substituent(s). Typical substituents include, but are not limited to, -M, -R<sup>60</sup>, -O<sup>-</sup>, =O, -OR<sup>60</sup>, -SR<sup>60</sup>, -S<sup>-</sup>, =S, -NR<sup>60</sup>R<sup>61</sup>, =NR<sup>60</sup>, -CF<sub>3</sub>, -CN, -OCN, -SCN, -NO, -NO<sub>2</sub>, =N<sub>2</sub>, -N<sub>3</sub>, -S(O)<sub>2</sub>O<sup>-</sup>, -S(O)<sub>2</sub>OH, -S(O)<sub>2</sub>R<sup>60</sup>, -OS(O<sub>2</sub>)O<sup>-</sup>, -OS(O)<sub>2</sub>R<sup>60</sup>, -P(O)(O<sup>-</sup>)<sub>2</sub>, -P(O)(OR<sup>60</sup>)(O<sup>-</sup>), -OP(O)(OR<sup>60</sup>)(OR<sup>61</sup>), -C(O)R<sup>60</sup>, -C(S)R<sup>60</sup>, -C(O)OR<sup>60</sup>, -C(O)NR<sup>60</sup>R<sup>61</sup>, -C(O)O<sup>-</sup>, -C(S)OR<sup>60</sup>, -NR<sup>62</sup>C(O)NR<sup>60</sup>R<sup>61</sup>, -NR<sup>62</sup>C(S)NR<sup>60</sup>R<sup>61</sup>, -NR<sup>62</sup>C(NR<sup>63</sup>)NR<sup>60</sup>R<sup>61</sup> and -C(NR<sup>62</sup>)NR<sup>60</sup>R<sup>61</sup> where M is independently a halogen; R<sup>60</sup>, R<sup>61</sup>, R<sup>62</sup> and R<sup>63</sup> are independently hydrogen, alkyl, substituted alkyl, alkoxy, substituted alkoxy, cycloalkyl, substituted cycloalkyl, cycloheteroalkyl, substituted cycloheteroalkyl, aryl, substituted aryl, heteroaryl or substituted heteroaryl, or optionally R<sup>60</sup> and R<sup>61</sup> together with the nitrogen atom to which they are bonded form a cycloheteroalkyl or substituted cycloheteroalkyl ring; and R<sup>64</sup> and R<sup>65</sup> are independently hydrogen, alkyl, substituted alkyl, aryl, cycloalkyl, substituted cycloalkyl, cycloheteroalkyl, substituted cycloheteroalkyl, aryl, substituted aryl, heteroaryl or substituted heteroaryl, or optionally R<sup>64</sup> and R<sup>65</sup> together with the nitrogen atom to which they are bonded form a cycloheteroalkyl or substituted cycloheteroalkyl ring. Preferably, substituents include -M, -R<sup>60</sup>, =O, -OR<sup>60</sup>, -SR<sup>60</sup>, -S<sup>-</sup>, =S, -NR<sup>60</sup>R<sup>61</sup>, =NR<sup>60</sup>, -CF<sub>3</sub>, -CN, -OCN, -SCN, -NO, -NO<sub>2</sub>, =N<sub>2</sub>, -N<sub>3</sub>, -S(O)<sub>2</sub>R<sup>60</sup>, -OS(O<sub>2</sub>)O<sup>-</sup>, -OS(O)<sub>2</sub>R<sup>60</sup>, -P(O)(O<sup>-</sup>)<sub>2</sub>,  
20  
25  
30

-P(O)(OR<sup>60</sup>)(O<sup>-</sup>), -OP(O)(OR<sup>60</sup>)(OR<sup>61</sup>), -C(O)R<sup>60</sup>, -C(S)R<sup>60</sup>, -C(O)OR<sup>60</sup>,  
 -C(O)NR<sup>60</sup>R<sup>61</sup>, -C(O)O<sup>-</sup>, -NR<sup>62</sup>C(O)NR<sup>60</sup>R<sup>61</sup>, more preferably, -M, -R<sup>60</sup>, =O, -OR<sup>60</sup>,  
 -SR<sup>60</sup>, -NR<sup>60</sup>R<sup>61</sup>, -CF<sub>3</sub>, -CN, -NO<sub>2</sub>, -S(O)<sub>2</sub>R<sup>60</sup>, -P(O)(OR<sup>60</sup>)(O<sup>-</sup>), -OP(O)(OR<sup>60</sup>)(OR<sup>61</sup>),  
 -C(O)R<sup>60</sup>, -C(O)OR<sup>60</sup>, -C(O)NR<sup>60</sup>R<sup>61</sup>, -C(O)O<sup>-</sup>, most preferably, -M, -R<sup>60</sup>, =O, -OR<sup>60</sup>,  
 5 -SR<sup>60</sup>, -NR<sup>60</sup>R<sup>61</sup>, -CF<sub>3</sub>, -CN, -NO<sub>2</sub>, -S(O)<sub>2</sub>R<sup>60</sup>, -OP(O)(OR<sup>60</sup>)(OR<sup>61</sup>), -C(O)R<sup>60</sup>,  
 -C(O)OR<sup>60</sup>, -C(O)O<sup>-</sup>, where R<sup>60</sup>, R<sup>61</sup> and R<sup>62</sup> are as defined above.

“Treating” or “treatment” of any disease or disorder refers, in one  
 embodiment, to ameliorating the disease or disorder (*i.e.*, arresting or reducing the  
 10 development of the disease or at least one of the clinical symptoms thereof). In  
 another embodiment “treating” or “treatment” refers to ameliorating at least one  
 physical parameter, which may not be discernible by the patient. In yet another  
 embodiment, “treating” or “treatment” refers to inhibiting the disease or disorder,  
 either physically, (*e.g.*, stabilization of a discernible symptom), physiologically, (*e.g.*,  
 15 stabilization of a physical parameter), or both. In yet another embodiment, “treating”  
 or “treatment” refers to delaying the onset of the disease or disorder.

“Therapeutically effective amount” means the amount of a compound that,  
 when administered to a patient for treating a disease, is sufficient to effect such  
 20 treatment for the disease. The “therapeutically effective amount” will vary depending  
 on the compound, the disease and its severity and the age, weight, *etc.*, of the patient  
 to be treated.

Reference will now be made in detail to preferred embodiments of the  
 25 invention. While the invention will be described in conjunction with the preferred  
 embodiments, it will be understood that it is not intended to limit the invention to  
 those preferred embodiments. To the contrary, it is intended to cover alternatives,  
 modifications, and equivalents as may be included within the spirit and scope of the  
 invention as defined by the appended claims.

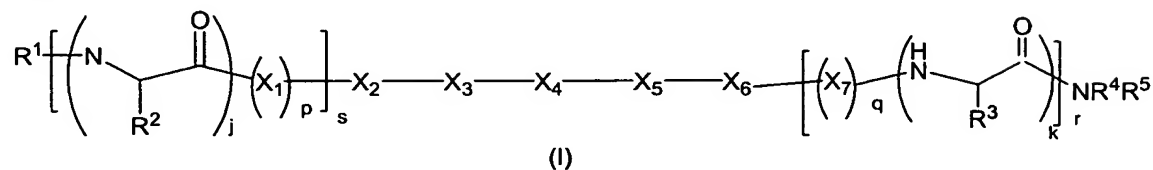
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#### 4.2 Compounds of the Invention

The present invention satisfies these and other needs by providing peptide  
 analogs of Ac-PHSCN-NH<sub>2</sub> which target tumor and endothelial cells and have anti-  
 tumor, anti-angiogenic and anti-metastatic activity, methods of making these

peptides, compositions thereof and methods of using these peptides and pharmaceutical compositions thereof to treat, prevent and detect diseases characterized by tumor growth, metastasis and angiogenesis. The peptide analogs may serve, *inter alia*, as carriers of radioactivity, PET-active compounds, toxins, fluorescent molecules and PEG molecules.

Accordingly, in one aspect, the present invention provides a compound of Formula (I):



or a pharmaceutically acceptable salt, solvate, hydrate or N-oxide thereof wherein:

j and k are independently 0 or 1;

p and q are independently an integer including and between 0 and 100;

r and s are independently 0 or 1;

R<sup>1</sup> is acyl, substituted acyl, acyl chelate, alkyl, substituted alkyl, cycloalkyl, substituted cycloalkyl, imino or substituted imino;

R<sup>2</sup> is C<sub>1</sub>-C<sub>6</sub> alkyl with at least one hydrogen atom replaced by a substituent selected from the group consisting of -NR<sup>6</sup>R<sup>7</sup>, -OR<sup>8</sup>, -CO<sub>2</sub>R<sup>9</sup>, -S(O)<sub>z</sub>R<sup>10</sup>, -P(OR<sup>11</sup>)OR<sup>12</sup>, aryl and substituted aryl;

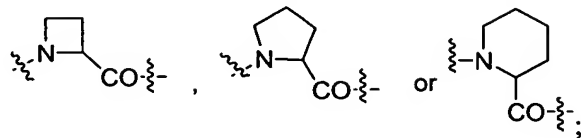
R<sup>6</sup>, R<sup>7</sup>, R<sup>8</sup>, R<sup>9</sup>, R<sup>10</sup>, R<sup>11</sup> and R<sup>12</sup> are independently selected from the group consisting of hydrogen, acyl, substituted acyl, acyl chelate, alkyl, substituted alkyl, cycloalkyl, substituted cycloalkyl, imino and substituted imino;

X<sub>1</sub> is -NH(C=C)<sub>g</sub>CO-, -NH(CH<sub>2</sub>)<sub>h</sub>CO- or -NHCH(CH<sub>3</sub>)CO-;



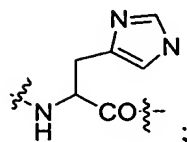
g and h are independently 1, 2, 3, 4, 5 or 6

X<sub>2</sub> is

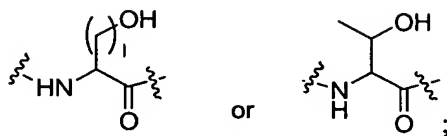


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X<sub>3</sub> is



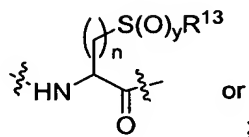
X<sub>4</sub> is



10

l is an integer from 1 to 4;

X<sub>5</sub> is



15

R<sup>13</sup> is hydrogen, alkyl, substituted alkyl, acyl, substituted acyl, arylalkyl, substituted arylalkyl, aryl, substituted aryl or -S(O)<sub>x</sub>R<sup>14</sup>;

n is an integer from 1 to 5;

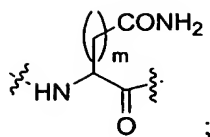
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R<sup>14</sup> is alkyl, substituted alkyl, acyl, substituted acyl, arylalkyl, substituted arylalkyl, aryl or substituted aryl;

y and x are independently 0, 1 or 2;

25

X<sub>6</sub> is



m is an integer from 1, 2, 3 or 4;

5

X<sub>7</sub> is -NH(C=C)<sub>d</sub>CO-, -NH(CH<sub>2</sub>)<sub>e</sub>CO- or -NHCH(CH<sub>3</sub>)CO-;

d and e are independently 1, 2, 3, 4, 5 or 6;

10

R<sup>3</sup> is C<sub>1</sub>-C<sub>6</sub> alkyl with at least one hydrogen atom replaced by a substituent selected from the group consisting of -NR<sup>15</sup>R<sup>16</sup>, -OR<sup>17</sup>, -CO<sub>2</sub>R<sup>18</sup>, -S(O)<sub>n</sub>R<sup>19</sup>, -P(OR<sup>20</sup>)OR<sup>21</sup>, aryl and substituted aryl;

15

R<sup>4</sup> and R<sup>5</sup> are independently hydrogen, alkyl or substituted alkyl; and

R<sup>15</sup>, R<sup>16</sup>, R<sup>17</sup>, R<sup>18</sup>, R<sup>19</sup>, R<sup>20</sup> and R<sup>21</sup> are independently selected from the group consisting of hydrogen, acyl, substituted acyl, acyl chelate, alkyl, substituted alkyl, cycloalkyl, substituted cycloalkyl, imino and substituted imino;

20

with the proviso that R<sup>1</sup> is not acetyl when R<sup>4</sup> and R<sup>5</sup> are hydrogen and r and s are 0.

In one embodiment, R<sup>1</sup> is not acetyl when R<sup>4</sup> and R<sup>5</sup> are hydrogen.

25

In one embodiment, p and q are independently an integer between and including 1 and 50. In another embodiment, p and q are independently an integer between and including 1 and 25. In still another embodiment, p and q are independently an integer between and including 1 and 10. In still another embodiment, p and q are independently an integer between and including 1 and 5. In still another embodiment, p and q are independently an integer between and including 1 and 3.

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In one embodiment, p and q are independently an integer between and including 0 and 50. In another embodiment, p and q are independently an integer between and including 0 and 25. In still another embodiment, p and q are

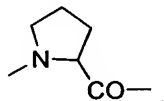
independently an integer between and including 0 and 10. In still another embodiment, p and q are independently an integer between and including 0 and 5. In still another embodiment, p and q are independently an integer between and including 0 and 3.

- 5 In still another embodiment, s is 0 and r is 1. In still another embodiment, s is 1 and r is 0. In still another embodiment, at least one of s and r is not 0.

In still another embodiment,  $R^1$  is acyl, substituted acyl, acyl chelate, imino or substituted imino. In still another embodiment,  $R^2$  is  $C_1$ - $C_6$  alkyl with at least one hydrogen atom replaced by a substituent selected from the group consisting of -  
 10  $NR^6R^7$ ,  $-OR^8$  and  $-CO_2R^9$ . In still another embodiment,  $R^6$ ,  $R^7$ ,  $R^8$ ,  $R^9$ ,  $R^{10}$ ,  $R^{11}$  and  $R^{12}$  are independently selected from the group consisting of hydrogen, acyl, substituted acyl, acyl chelate, imino or substituted imino.

In still another embodiment,  $X_1$  is  $-NHCH_2CO-$  or  $-NHCH(CH_3)CO-$ . In still another embodiment,  $X_2$  is

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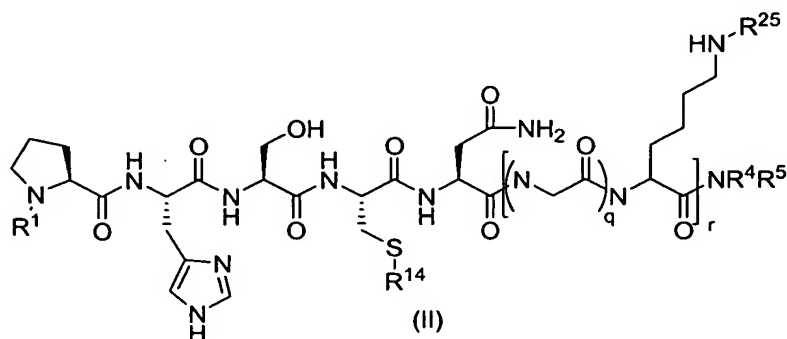


In still another embodiment, l is 1. In still another embodiment, n is 1 or 2. In still another embodiment, m is 1 or 2.

In still another embodiment,  $R^3$  is  $C_1$ - $C_6$  alkyl with at least one hydrogen atom replaced by a substituent selected from the group consisting of  $-NR^{15}R^{16}$ ,  $-OR^{17}$  and  $-CO_2R^{18}$ . In still another embodiment,  $R^{15}$ ,  $R^{16}$ ,  $R^{17}$ ,  $R^{18}$ ,  $R^{19}$ ,  $R^{20}$  and  $R^{21}$  are  
 20 independently selected from the group consisting of hydrogen, acyl, substituted acyl, acyl chelate, imino or substituted imino.

In one embodiment, compounds of the invention has the structure of Formula (II):

25



wherein:

R<sup>1</sup> is acyl, substituted acyl, acyl chelate, alkyl, substituted alkyl, imino or substituted imino;

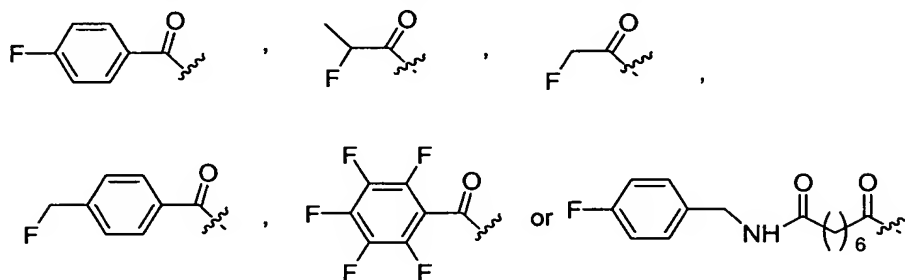
5 R<sup>4</sup> is hydrogen;

R<sup>5</sup> is hydrogen, alkyl or substituted alkyl; and

10 R<sup>25</sup> is hydrogen, acyl, substituted acyl, acyl chelate, alkyl, substituted alkyl, imino or substituted imino; and

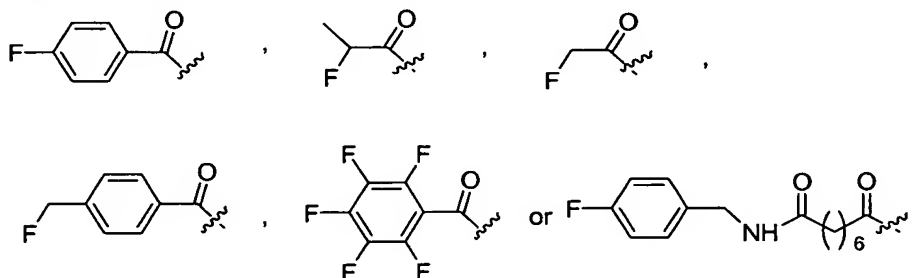
q is 1, 2, 3, 4 or 5.

In one embodiment of a compound of Formulae (I) and (II), R<sup>1</sup> is



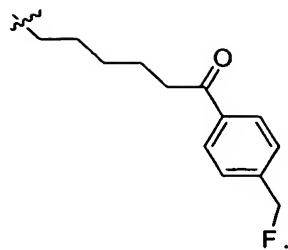
15 and R<sup>14</sup> is hydrogen, methyl or acetyl. In one specific embodiment, r = 0 and R<sup>4</sup> and R<sup>5</sup> are hydrogen. In another specific embodiment, r = 1, q is 2, R<sup>4</sup>, R<sup>5</sup> and R<sup>25</sup> are hydrogen.

In another embodiment of a compound of Formulae (I) and (II), r = 1, q is 2, R<sup>14</sup> is hydrogen, methyl or acetyl and R<sup>25</sup> is

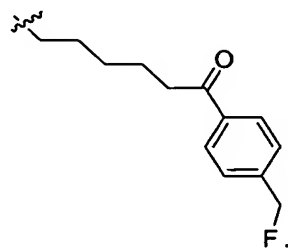


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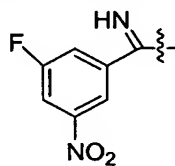
In another embodiment of a compound of Formulae (I) and (II), r = 0, R<sup>1</sup> is acetyl, R<sup>4</sup> is hydrogen, R<sup>14</sup> is hydrogen, methyl or acetyl and R<sup>5</sup> is



In still another embodiment of a compound of Formulae (I) and (II),  $r = 1$ ,  $q$  is 2,  $R^1$  is acetyl,  $R^4$  and  $R^{25}$  are hydrogen,  $R^{14}$  is hydrogen, methyl or acetyl and  $R^5$  is



5 In still another embodiment of a compound of Formulae (I) and (II),  $r = 0$ ,  $R^4$  and  $R^5$  are hydrogen,  $R^{14}$  is hydrogen, methyl or acetyl and  $R^1$  is

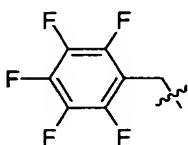
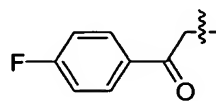
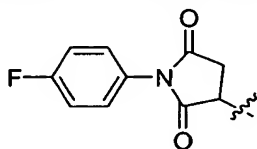


In still another embodiment of a compound of Formulae (I) and (II),  $r = 1$ ,  $q = 2$ ,  $R^1 =$  acetyl,  $R^4$  and  $R^5$  are hydrogen,  $R^{14}$  is hydrogen, methyl or acetyl and  $R^{25}$  is

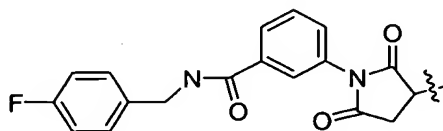


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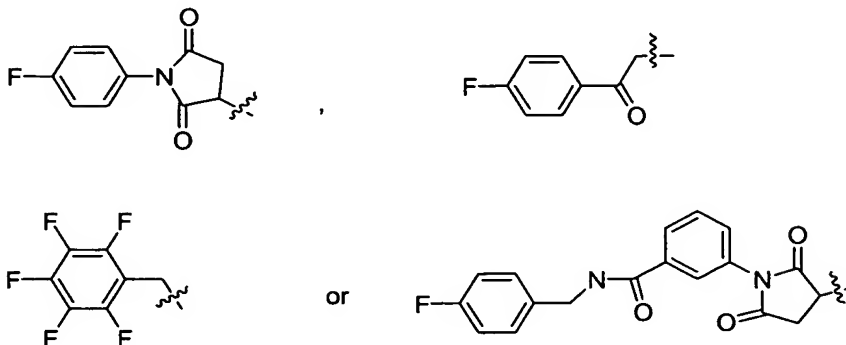
In still another embodiment of a compound of Formulae (I) and (II),  $r = 0$ ,  $R^4$  and  $R^5$  are hydrogen,  $R^{14}$  is hydrogen, methyl or acetyl and  $R^1$  is



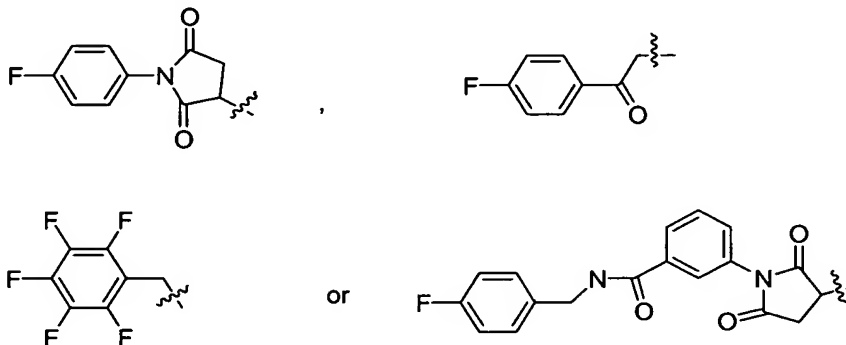
or



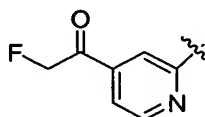
In still another embodiment of a compound of Formulae (I) and (II),  $r = 1$ ,  $q = 2$ ,  $R^4$ ,  $R^5$  and  $R^{25}$  are hydrogen,  $R^{14}$  is hydrogen, methyl or acetyl and  $R^1$  is



5 In still another embodiment of a compound of Formulae (I) and (II),  $r = 1$ ,  $q = 2$ ,  $R^1$  is acetyl,  $R^4$  and  $R^5$  are hydrogen,  $R^{14}$  is hydrogen, methyl or acetyl and  $R^{25}$  is

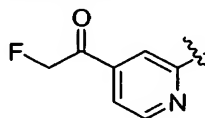


In still another embodiment of a compound of Formulae (I) and (II),  $r = 0$ ,  $R^4$  and  $R^5$  are hydrogen,  $R^{14}$  is hydrogen, methyl or acetyl and  $R^1$  is

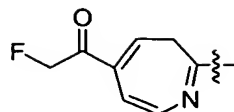


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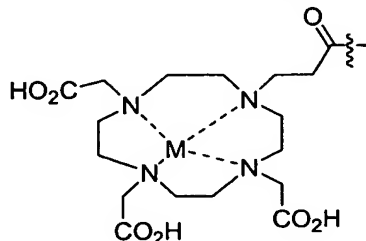
In still another embodiment of a compound of Formulae (I) and (II),  $r = 1$ ,  $q = 2$ ,  $R^4$ ,  $R^5$  and  $R^{25}$  are hydrogen,  $R^{14}$  is hydrogen, methyl or acetyl and  $R^1$  is



15 In still another embodiment of a compound of Formulae (I) and (II),  $r = 1$ ,  $q = 2$ ,  $R^1$  is acetyl,  $R^4$  and  $R^5$  are hydrogen,  $R^{14}$  is hydrogen, methyl or acetyl and  $R^{25}$  is

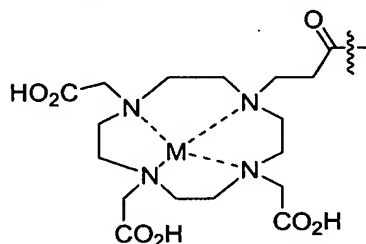


In still another embodiment of a compound of Formulae (I) and (II),  $r = 0$ ,  $R^4$  and  $R^5$  are hydrogen,  $R^{14}$  is hydrogen, methyl or acetyl and  $R^1$  is



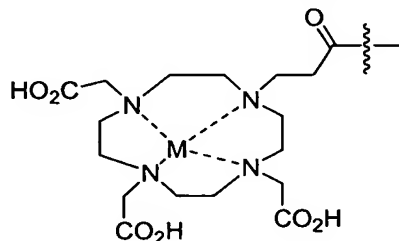
5 and M is Cu, Ga,  $^{111}\text{In}$  or  $^{90}\text{Y}$ .

In still another embodiment of a compound of Formulae (I) and (II),  $r = 1$ ,  $q = 2$ ,  $R^4$ ,  $R^5$  and  $R^{25}$  are hydrogen,  $R^{14}$  is hydrogen, methyl or acetyl,  $R^1$  is



and M is Cu, Ga,  $^{111}\text{In}$  or  $^{90}\text{Y}$ .

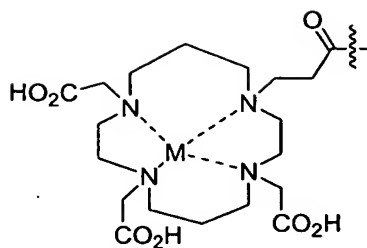
10 In still another embodiment of a compound of Formulae (I) and (II),  $r = 1$ ,  $q = 2$ ,  $R^1$  is acetyl,  $R^4$  and  $R^5$  are hydrogen,  $R^{14}$  is hydrogen, methyl or acetyl,  $R^{25}$  is



and M is Cu, Ga,  $^{111}\text{In}$  or  $^{90}\text{Y}$ .

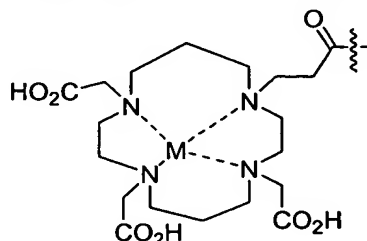
In still another embodiment of a compound of Formulae (I) and (II),  $r = 0$ ,  $R^4$  and  $R^5$  are hydrogen,  $R^{14}$  is hydrogen, methyl or acetyl and  $R^1$  is

15



and M is Cu.

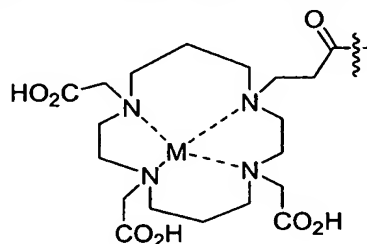
In still another embodiment of a compound of Formulae (I) and (II),  $r = 1$ ,  $q = 2$ ,  $R^4$ ,  $R^5$  and  $R^{25}$  are hydrogen,  $R^{14}$  is hydrogen, methyl or acetyl,  $R^1$  is



5

and M is Cu.

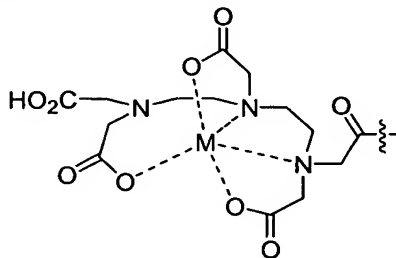
In still another embodiment of a compound of Formulae (I) and (II),  $r = 1$ ,  $q = 2$ ,  $R^1$  is acetyl,  $R^4$  and  $R^5$  are hydrogen,  $R^{14}$  is hydrogen, methyl or acetyl,  $R^{25}$  is



10

and M is Cu.

In still another embodiment of a compound of Formulae (I) and (II),  $r = 0$ ,  $R^4$  and  $R^5$  are hydrogen,  $R^{14}$  is hydrogen, methyl or acetyl and  $R^1$  is

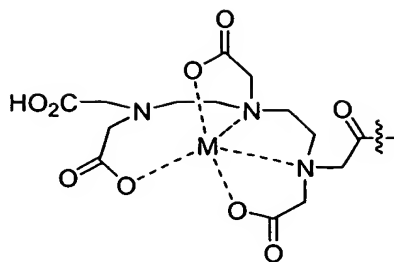


and M is  $^{111}\text{In}$ ,  $^{90}\text{Y}$  or Ga.

15

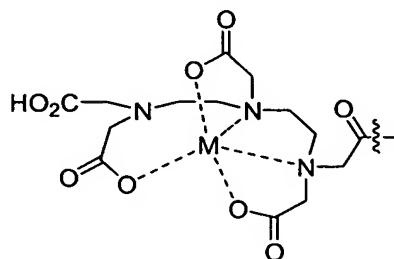
In still another embodiment of a compound of Formulae (I) and (II),  $r = 1$ ,  $q = 2$ ,  $R^4$ ,  $R^5$  and  $R^{25}$  are hydrogen,  $R^{14}$  is hydrogen, methyl or acetyl,  $R^1$  is





and M is  $^{111}\text{In}$ ,  $^{90}\text{Y}$  or Ga.

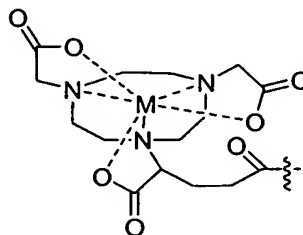
In still another embodiment of a compound of Formulae (I) and (II),  $r = 1$ ,  $q = 2$ ,  $R^1$  is acetyl,  $R^4$  and  $R^5$  are hydrogen,  $R^{14}$  is hydrogen, methyl or acetyl,  $R^{25}$  is



5

and M is  $^{111}\text{In}$ ,  $^{90}\text{Y}$  or Ga.

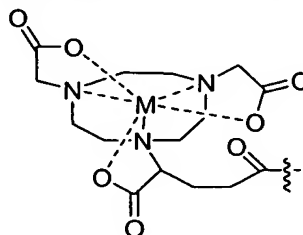
In still another embodiment of a compound of Formulae (I) and (II),  $r = 0$ ,  $R^4$  and  $R^5$  are hydrogen,  $R^{14}$  is hydrogen, methyl or acetyl,  $R^1$  is



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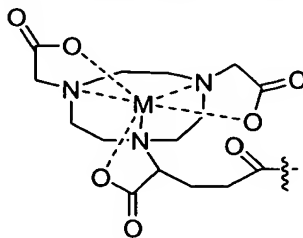
and M is  $^{111}\text{In}$ ,  $^{67}\text{Ga}$  or  $^{68}\text{Ga}$ .

In still another embodiment of a compound of Formulae (I) and (II),  $r = 1$ ,  $q = 2$ ,  $R^4$ ,  $R^5$  and  $R^{25}$  are hydrogen,  $R^{14}$  is hydrogen, methyl or acetyl,  $R^1$  is



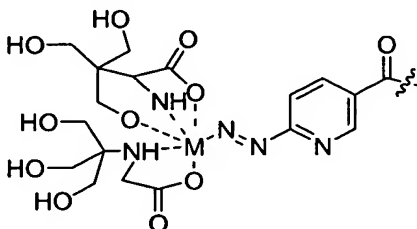
15 and M is  $^{111}\text{In}$ ,  $^{67}\text{Ga}$  or  $^{68}\text{Ga}$ .

In still another embodiment of a compound of Formulae (I) and (II),  $r = 1$ ,  $q = 2$ ,  $R^1$  is acetyl,  $R^4$  and  $R^5$  are hydrogen,  $R^{14}$  is hydrogen, methyl or acetyl,  $R^{25}$  is



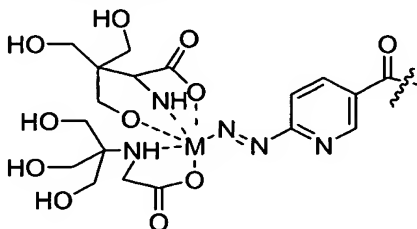
and M is  $^{111}\text{In}$ ,  $^{67}\text{Ga}$  or  $^{68}\text{Ga}$ .

- 5 In still another embodiment of a compound of Formulae (I) and (II), wherein  $r = 0$ ,  $R^4$  and  $R^5$  are hydrogen,  $R^{14}$  is hydrogen, methyl or acetyl,  $R^1$  is



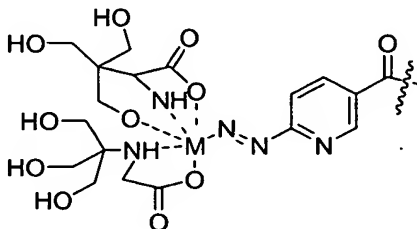
and M is Tc.

- 10 In still another embodiment of a compound of Formulae (I) and (II),  $r = 1$ ,  $q = 2$ ,  $R^4$ ,  $R^5$  and  $R^{25}$  are hydrogen,  $R^{14}$  is hydrogen, methyl or acetyl,  $R^1$  is



and M is Tc.

- 15 In still another embodiment of a compound of Formulae (I) and (II),  $r = 1$ ,  $q = 2$ ,  $R^1$  is acetyl,  $R^4$  and  $R^5$  are hydrogen,  $R^{14}$  is hydrogen, methyl or acetyl,  $R^{25}$  is



and M is Tc.

compounds to be tested are implanted into tiny incisions made in the cornea 1.0-1.5 mm from the limbus. Neovascularization is assessed at 5 and 7 days after implantation. On day 7, animals are anesthetized and infused with a dye such as colloidal carbon to stain the vessels. The animals are then euthanized, the corneas  
5 fixed with formalin, and the corneas flattened and photographed to assess the degree of neovascularization. Neovessels may be quantitated by imaging the total vessel area or length or simply by counting vessels.

#### 4.4.8 Matrigel® Plug Assay

10 This assay is performed essentially as described by Passaniti *et al.*, 1992, *Lab Invest.* 67:519-528. Ice-cold Matrigel® (*e.g.*, 500  $\mu$ L) (Collaborative Biomedical Products, Inc., Bedford, MA) is mixed with heparin (*e.g.*, 50  $\mu$ g/ml), FGF-2 (*e.g.*, 400 ng/ml) and the compound to be tested. In some assays, bFGF may be substituted with  
15 tumor cells as the angiogenic stimulus. The Matrigel® mixture is injected subcutaneously into 4-8 week-old athymic nude mice at sites near the abdominal midline, preferably 3 injections per mouse. The injected Matrigel® forms a palpable solid gel. Injection sites are chosen such that each animal receives a positive control plug (such as FGF-2 + heparin), a negative control plug (*e.g.*, buffer + heparin) and a  
20 plug that includes the compound being tested for its effect on angiogenesis, *e.g.*, (FGF-2 + heparin + compound). All treatments are preferably run in triplicate. Animals are sacrificed by cervical dislocation at about 7 days post injection or another time that may be optimal for observing angiogenesis. The mouse skin is detached along the abdominal midline, and the Matrigel® plugs are recovered and scanned immediately at high resolution. Plugs are then dispersed in water and incubated at  
25 37°C overnight. Hemoglobin (Hb) levels are determined using Drabkin's solution (*e.g.*, obtained from Sigma) according to the manufacturers' instructions. The amount of Hb in the plug is an indirect measure of angiogenesis as it reflects the amount of blood in the sample. In addition, or alternatively, animals may be injected prior to sacrifice with a 0.1 ml buffer (preferably PBS) containing a high molecular weight  
30 dextran to which is conjugated a fluorophore. The amount of fluorescence in the dispersed plug, determined fluorimetrically, also serves as a measure of angiogenesis in the plug. Staining with mAb anti-CD31 (CD31 is "platelet-endothelial cell

adhesion molecule or PECAM”) may also be used to confirm neovessel formation and microvessel density in the plugs.

#### **4.4.9 Chick Chorioallantoic Membrane (CAM) Angiogenesis Assay**

5 This assay is performed essentially as described by Nguyen *et al.*,  
*Microvascular Res.* 1994, 47:31-40. A mesh containing either angiogenic factors  
(bFGF) or tumor cells plus inhibitors is placed onto the CAM of an 8-day old chick  
embryo and the CAM observed for 3-9 days after implantation of the sample.  
Angiogenesis is quantitated by determining the percentage of squares in the mesh  
10 which contain blood vessels.

#### **4.4.10 In Vivo Assessment of Angiogenesis Inhibition and Anti-Tumor Effects Using the Matrigel® Plug Assay with Tumor Cells**

In this assay, tumor cells, for example  $1-5 \times 10^6$  cells of the 3LL Lewis lung  
15 carcinoma or the rat prostate cell line MatLyLu, are mixed with Matrigel® and then  
injected into the flank of a mouse following the protocol described in Sec. B., above.  
A mass of tumor cells and a powerful angiogenic response can be observed in the  
plugs after about 5 to 7 days. The anti-tumor and anti-angiogenic action of a  
compound in an actual tumor environment can be evaluated by including it in the  
20 plug. Measurement is then made of tumor weight, Hb levels or fluorescence levels  
(of a dextran-fluorophore conjugate injected prior to sacrifice). To measure Hb or  
fluorescence, the plugs are first homogenize with a tissue homogenizer.

#### **4.4.11 Xenograft Model of Subcutaneous (s.c.) Tumor Growth**

25 Nude mice are inoculated with MDA-MB-231 cells (human breast carcinoma)  
and Matrigel® ( $1 \times 10^6$  cells in 0.2mL) s.c. in the right flank of the animals. The  
tumors are staged to 200 mm<sup>3</sup> and then treatment with a test composition is initiated  
(100µg/animal/day given q.d. IP). Tumor volumes are obtained every other day and  
the animals are sacrificed after 2 weeks of treatment. The tumors are excised,  
30 weighed and paraffin embedded. Histological sections of the tumors are analyzed by  
H and E, anti-CD31, Ki-67, TUNEL, and CD68 staining.

#### 4.4.12 Xenograft Model of Metastasis

The compounds of the invention are also tested for inhibition of late metastasis using an experimental metastasis model (Crowley *et al.*, *Proc. Natl. Acad. Sci. USA* 5 1993, 90 5021-5025). Late metastasis involves the steps of attachment and extravasation of tumor cells, local invasion, seeding, proliferation and angiogenesis. Human prostatic carcinoma cells (PC-3) transfected with a reporter gene, preferably the green fluorescent protein (GFP) gene, but as an alternative with a gene encoding the enzymes chloramphenicol acetyl-transferase (CAT), luciferase or LacZ, are 10 inoculated into nude mice. This approach permits utilization of either of these markers (fluorescence detection of GFP or histochemical colorimetric detection of enzymatic activity) to follow the fate of these cells. Cells are injected, preferably iv, and metastases identified after about 14 days, particularly in the lungs but also in regional lymph nodes, femurs and brain. This mimics the organ tropism of naturally 15 occurring metastases in human prostate cancer. For example, GFP-expressing PC-3 cells ( $1 \times 10^6$  cells per mouse) are injected iv into the tail veins of nude (*nu/nu*) mice. Animals are treated with a test composition at 100µg/animal/day given q.d. IP. Single metastatic cells and foci are visualized and quantitated by fluorescence microscopy or light microscopic histochemistry or by grinding the tissue and quantitative 20 colorimetric assay of the detectable label.

#### 4.4.13. Inhibition of Spontaneous Metastasis *In Vivo* by PHSCN and Functional Derivatives

The rat syngeneic breast cancer system employs Mat BIII rat breast cancer 25 cells (Xing *et al.*, *Int. J. Cancer* 1996, 67:423-429). Tumor cells, for example, about  $10^6$  suspended in 0.1 mL PBS, are inoculated into the mammary fat pads of female Fisher rats. At the time of inoculation, a 14-day Alza osmotic mini-pump is implanted intraperitoneally to dispense the test compound. The compound is dissolved in PBS (*e.g.*, 200 mM stock), sterile filtered and placed in the minipump to 30 achieve a release rate of about 4 mg/kg/day. Control animals receive vehicle (PBS) alone or a vehicle control peptide in the minipump. Animals are sacrificed at about day 14. In the rats treated with the compounds of the present invention, significant reductions in the size of the primary tumor and in the number of metastases in the

spleen, lungs, liver, kidney and lymph nodes (enumerated as discrete foci) may be observed. Histological and immunohistochemical analysis reveal increased necrosis and signs of apoptosis in tumors in treated animals. Large necrotic areas are seen in tumor regions lacking neovascularization. Human or rabbit PHSCN and their derivatives to which  $^{131}\text{I}$  is conjugated (either 1 or 2 I atoms per molecule of peptide) are effective radiotherapeutics and are found to be at least two-fold more potent than the unconjugated polypeptides. In contrast, treatment with control peptides fails to cause a significant change in tumor size or metastasis.

#### **4.4.14. 3LL Lewis Lung Carcinoma: Primary Tumor Growth**

This tumor line arose spontaneously as carcinoma of the lung in a C57BL/6 mouse (Malave *et al.*, *J. Nat'l. Canc. Inst.* **1979**, 62:83-88). It is propagated by passage in C57BL/6 mice by subcutaneous (sc) inoculation and is tested in semiallogeneic C57BL/6 x DBA/2 F<sub>1</sub> mice or in allogeneic C3H mice. Typically six animals per group for subcutaneously (sc) implant, or ten for intramuscular (im) implant are used. Tumor may be implanted sc as a 2-4 mm fragment, or im or sc as an inoculum of suspended cells of about  $0.5\text{-}2 \times 10^6$ -cells. Treatment begins 24 hours after implant or is delayed until a tumor of specified size (usually approximately 400 mg) can be palpated. The test compound is administered ip daily for 11 days. Animals are followed by weighing, palpation, and measurement of tumor size. Typical tumor weight in untreated control recipients on day 12 after im inoculation is 500-2500 mg. Typical median survival time is 18-28 days. A positive control compound, for example cyclophosphamide at 20 mg/kg/injection per day on days 1-11 is used. Results computed include mean animal weight, tumor size, tumor weight, survival time. For confirmed therapeutic activity, the test composition should be tested in two multi-dose assays.

#### **4.4.15 3LL Lewis Lung Carcinoma: Primary Growth and Metastasis**

##### **Model**

This assay is well known in the art (Gorelik *et al.*, *J. Nat'l. Canc. Inst.* **1980**, 65:1257-1264; Gorelik *et al.*, *Rec. Results Canc. Res.* **1980**, 75:20-28; Isakov *et al.*, *Invasion Metas.* 2:12-32 (1982); Talmadge *et al.*, *J. Nat'l. Canc. Inst.* **1982**, 69:975-980; Hilgard *et al.*, *Br. J. Cancer* **1977**, 35:78-86). Test mice are male C57BL/6 mice, 2-3 months old. Following sc, im, or intra-footpad implantation, this

tumor produces metastases, preferentially in the lungs. With some lines of the tumor, the primary tumor exerts anti-metastatic effects and must first be excised before study of the metastatic phase (see also U.S. Patent No. 5,639,725).

Single-cell suspensions are prepared from solid tumors by treating minced tumor  
5 tissue with a solution of 0.3% trypsin. Cells are washed 3 times with PBS (pH 7.4) and suspended in PBS. Viability of the 3LL cells prepared in this way is generally about 95-99% (by trypan blue dye exclusion). Viable tumor cells ( $3 \times 10^4$  -  $5 \times 10^6$ ) suspended in 0.05 ml PBS are injected subcutaneously, either in the dorsal region or into one hind foot pad of C57BL/6 mice. Visible tumors appear after 3-4 days after  
10 dorsal sc injection of  $10^6$  cells. The day of tumor appearance and the diameters of established tumors are measured by caliper every two days. The treatment is given as one to five doses of peptide or derivative, per week. In another embodiment, the peptide is delivered by osmotic minipump.

In experiments involving tumor excision of dorsal tumors, when tumors reach  
15 about 1500 mm<sup>3</sup> in size, mice are randomized into two groups: (1) primary tumor is completely excised; or (2) sham surgery is performed and the tumor is left intact. Although tumors from 500-3000 mm<sup>3</sup> inhibit growth of metastases, 1500 mm<sup>3</sup> is the largest size primary tumor that can be safely resected with high survival and without local regrowth. After 21 days, all mice are sacrificed and autopsied.

20 Lungs are removed and weighed. Lungs are fixed in Bouin's solution and the number of visible metastases is recorded. The diameters of the metastases are also measured using a binocular stereoscope equipped with a micrometer-containing ocular under 8X magnification. On the basis of the recorded diameters, it is possible to calculate the volume of each metastasis. To determine the total volume of metastases  
25 per lung, the mean number of visible metastases is multiplied by the mean volume of metastases. To further determine metastatic growth, it is possible to measure incorporation of <sup>125</sup>IdUrd into lung cells (Thakur *et al.*, *J. Lab. Clin. Med.* 1977, 89:217-228). Ten days following tumor amputation, 25 µg of fluorodeoxyuridine is inoculated into the peritoneums of tumor-bearing (and, if used, tumor-resected mice).  
30 After 30 min, mice are given 1 µCi of <sup>125</sup>IdUrd (iododeoxyuridine). One day later, lungs and spleens are removed and weighed, and a degree of <sup>125</sup>IdUrd incorporation is measured using a gamma counter.

In mice with footpad tumors, when tumors reach about 8-10 mm in diameter, mice are randomized into two groups: (1) legs with tumors are amputated after ligation above the knee joints; or (2) mice are left intact as nonamputated tumor-bearing controls. (Amputation of a tumor-free leg in a tumor-bearing mouse  
5 has no known effect on subsequent metastasis, ruling out possible effects of anesthesia, stress or surgery). Mice are killed 10-14 days after amputation. Metastases are evaluated as described above.

Statistics: Values representing the incidence of metastases and their growth in the lungs of tumor-bearing mice are not normally distributed. Therefore,  
10 non-parametric statistics such as the Mann-Whitney U-Test may be used for analysis. Study of this model by Gorelik *et al.* (1980, *supra*) showed that the size of the tumor cell inoculum determined the extent of metastatic growth. The rate of metastasis in the lungs of operated mice was different from primary tumor-bearing mice. Thus in the lungs of mice in which the primary tumor had been induced by inoculation of  
15 larger doses of 3LL cells ( $1-5 \times 10^6$ ) followed by surgical removal, the number of metastases was lower than that in nonoperated tumor-bearing mice, though the volume of metastases was higher than in the nonoperated controls. Using  $^{125}\text{I}$ dUrd incorporation as a measure of lung metastasis, no significant differences were found between the lungs of tumor-excised mice and tumor-bearing mice originally  
20 inoculated with  $10^6$  3LL cells. Amputation of tumors produced following inoculation of  $10^5$  tumor cells dramatically accelerated metastatic growth. These results were in accord with the survival of mice after excision of local tumors. The phenomenon of acceleration of metastatic growth following excision of local tumors had been repeatedly observed (for example, see U. S. Patent No. 5,639,725). These  
25 observations have implications for the prognosis of patients who undergo cancer surgery.

#### 4.4.15 Competition Binding Assay with DU145 Prostate Cancer Cells.

30 These can be carried at 6 concentrations of peptide to determine an  $\text{IC}_{50}$ . This assay is a surrogate marker of biological activity. Briefly, DU145 cell are harvested by brief incubation with trypsin, the peptide derivative to be tested (competitor) and standard are added and the suspension is agitated at 4 °C for 2 hours. The cells are pelleted, supernatant aspirated, PBS added and the process is repeated three times



(washes). HRP-Streptavidin is added, allowed to bind followed by a second series of wash steps. The appropriate HRP substrate is added and color is developed within the previously defined linear part of the reaction.

5                   **4.4.16 Human Endothelial Cell (HUVEC) Proliferation Assay.**

The assay is carried out in a 96 well assay and takes place for 48 hours. Briefly, HUVECs (Cascade Biologicals) are cultured overnight in M200 containing 2% FBS. The following day, 3,000 cells are plated in each well of a gelatin-coated 96-well plate. The cells are allowed to adhere and spread for 4-6 hours, at which time  
10 the medium is replaced with fresh medium containing 2% FBS, 1 ng/mL FGF-2, and varying concentrations of specific test compounds. Cells are then cultured for an additional 48 hours, and relative cell numbers in each well are determined using the Cell Titer Aqueous Cell Proliferation Assay (Promega). The result will be an IC<sub>50</sub> that can be compared to a standard compound. This is an assay that reflects biological  
15 function of the peptide derivatives against their target cells.

**4.4.17 Dosing Range and Kinetics**

Four doses of peptide (2, 4, 8 and 10 µCi for biodistribution and 10, 50, 100, and 200 µCi for Gamma scintigraphy) are injected *via* tail vein into tumor (B16F10  
20 melanoma cells) bearing (200-300 mm<sup>3</sup>) C57Bl/6 mice. The lowest and highest dosages of peptides are tested first and second. Thus, it is possible that the dosage of peptide derivative to be tested may be modified based on the initial results obtained.

(Group I, Dosimetry 3 animals/dose) Animals will be sacrificed after 2 hours after dosing with <sup>125</sup>I labeled peptides. Tumors (theoretical high counts) and hearts  
25 (theoretical low counts) will then be removed and radioactivity counted.

(Group 2 Gamma scintigraphy 3 animals/dose) Animals are anesthetized and imaged (gamma scintigraphy) 2 hours after injection of <sup>125</sup>I radiolabeled peptide. A peptide dosage is chosen based on these results.

30                   **4.4.18 Imaging Feasibility**

<sup>125</sup>I labeled peptide at a dose determined by the method of section 4.4.17 will be injected *via* tail vein into tumor bearing mice (200-300 mm<sup>3</sup> tumor). Three mice per time point will be anesthetized and imaged at 0.5 h, 1 h, 2 h, 4 h, 6 h, 24 h, and 48

h which will determine the optimal time for imaging. As a control, unlabelled peptide at a 100-fold molar excess will be co-injected with the  $^{125}\text{I}$  labeled peptide to demonstrate equivalency of both species.

5

#### **4.5 Recombinant DNA Methods**

General methods of molecular biology have been amply described in the art (Sambrook, *et al.*, *Molecular Cloning: A Laboratory Manual*, 2nd (or later) Edition, Cold Spring Harbor Press, Cold Spring Harbor, NY, 1989; Ausube *et al.*, *Current Protocols in Molecular Biology*, Vol. 2, Wiley-Interscience, New York, (current edition); Kriegler, *Gene Transfer and Expression: A Laboratory Manual* (1990); Glover, DM, editor, *DNA Cloning: A Practical Approach*, vol. I & II, IRL Press, 1985; Alberts *et al.*, *Molecular Biology of the Cell*, 2nd (or later) Ed., Garland Publishing, Inc., New York, NY (1989); Watson *et al.*, *Recombinant DNA*, 2nd (or later) Ed., Scientific American Books, New York, 1992; and Old *et al.*, *Principles of Gene Manipulation: An Introduction to Genetic Engineering*, 2nd (or later) Ed., University of California Press, Berkeley, CA (1981)).

Unless otherwise indicated, a particular nucleic acid sequence is intended to encompass conservative substitution variants thereof (*e.g.*, degenerate codon substitutions) and a complementary sequence. The term “nucleic acid” is synonymous with “polynucleotide” and is intended to include a gene, a cDNA molecule, an mRNA molecule, as well as a fragment of any of these such as an oligonucleotide, and further, equivalents thereof (explained more fully below). Sizes of nucleic acids are stated either as kilobases (kb) or base pairs (bp). These are estimates derived from agarose or polyacrylamide gel electrophoresis (PAGE), from nucleic acid sequences which are determined by the user or published. Protein size is stated as molecular mass in kilodaltons (kDa) or as length (number of amino acid residues). Protein size is estimated from PAGE, from sequencing, from presumptive amino acid sequences based on the coding nucleic acid sequence or from published amino acid sequences.

30

Specifically, DNA molecules encoding the amino acid sequence corresponding to the peptide multimers of the present invention, or active variants thereof, can be synthesized by the polymerase chain reaction (PCR) (see, for example, U. S. Patent No. 4,683,202) using primers derived the sequence of the protein disclosed herein. These cDNA sequences can then be assembled into a eukaryotic or

prokaryotic expression vector and the resulting vector can be used to direct the synthesis of the fusion polypeptide or its fragment or derivative by appropriate host cells, for example COS or CHO cells.

The term "nucleic acid" as used herein is intended to include such fragments or equivalents. The nucleic acid sequences of this invention can be DNA or RNA. Prokaryotic or eukaryotic host cells transformed or transfected to express the multimers are within the scope of the invention. For example, the peptide multimer may be expressed in bacterial cells such as *E. coli*, insect cells (baculovirus), yeast, or mammalian cells such as Chinese hamster ovary cells (CHO) or human cells (which are preferred for human therapeutic use of the transfected cells). Other suitable host are known to those skilled in the art. Expression in eukaryotic cells leads to partial or complete glycosylation and/or formation of relevant inter- or intra-chain disulfide bonds of the recombinant polypeptide. Examples of vectors for expression in yeast *S. cerevisiae* include pYepSec1 (Baldari *et al.*, 1987, *EMBO J.* 6:229-234), pMFa (Kurjan *et al.* 1982 *Cell* 30:933-943), pJRY88 (Schultz *et al.*, 1987, *Gene* 54:113-123), and pYES2 (Invitrogen Corporation, San Diego, Calif.). Baculovirus vectors available for expression of proteins in cultured insect cells (SF 9 cells) include the pAc series (Smith *et al.*, 1983, *Mol. Cell Biol.* 3:2156-2165) and the pVL series (Lucklow *et al.*, (1989) *Virology* 170:31-39). Generally, COS cells (Gluzman 1981 *Cell* 23:175-182) are used in conjunction with such vectors as pCDM 8 (Aruffo *et al.*, *supra*, for transient amplification/expression in mammalian cells, while CHO (*dhfr*-negative CHO) cells are used with vectors such as pMT2PC (Kaufman *et al.*, 1987, *EMBO J.* 6:187-195) for stable amplification/expression in mammalian cells. The NS0 myeloma cell line (a glutamine synthetase expression system.) is available from Celltech Ltd.

Construction of suitable vectors containing the desired coding and control sequences employs standard ligation and restriction techniques which are well understood in the art. Isolated plasmids, DNA sequences, or synthesized oligonucleotides are cleaved, tailored, and re-ligated in the form desired. The DNA sequences which form the vectors are available from a number of sources. Backbone vectors and control systems are generally found on available "host" vectors which are used for the bulk of the sequences in construction. For the pertinent coding sequence, initial construction may be, and usually is, a matter of retrieving the appropriate sequences from cDNA or genomic DNA libraries. However, once the sequence is

disclosed it is possible to synthesize the entire gene sequence *in vitro* starting from the individual nucleotide derivatives. The entire gene sequence for genes of sizeable length, *e.g.*, 500-1000 bp may be prepared by synthesizing individual overlapping complementary oligonucleotides and filling in single stranded nonoverlapping portions using DNA polymerase in the presence of the deoxyribonucleotide triphosphates. This approach has been used successfully in the construction of several genes of known sequence. See, for example, Edge, *Nature* **1981**, 292:756; Nambair *et al.*, *Science* **1984**, 223:1299; and Jay, *J. Biol. Chem.* **1984**, 259:6311.

Synthetic oligonucleotides are prepared by either the phosphotriester method as described by references cited above or the phosphoramidite method as described by Beaucage *et al.*, *Tetrahedron Lett.* **1981**, 22:1859; and Matteucci *et al.*, *J. Am. Chem. Soc.* **1981**, 103:3185 and can be prepared using commercially available automated oligonucleotide synthesizers. Kinase treatment of single strands prior to annealing or for labeling is achieved using well-known methods.

Once the components of the desired vectors are thus available, they can be excised and ligated using standard restriction and ligation procedures. Site-specific DNA cleavage is performed by treating with the suitable restriction enzyme (or enzymes) under conditions which are generally understood in the art, and the particulars of which are specified by the manufacturer of these commercially available restriction enzymes. See, *e.g.*, New England Biolabs, Product Catalog. If desired, size separation of the cleaved fragments may be performed by polyacrylamide gel or agarose gel electrophoresis using standard techniques. A general description of size separations is found in *Meth. Enzymol.* (1980) 65:499-560. Any of a number of methods are used to introduce mutations into the coding sequence to generate variants if these are to be produced recombinantly. These mutations include simple deletions or insertions, systematic deletions, insertions or substitutions of clusters of bases or substitutions of single bases. Modifications of the DNA sequence are created by site-directed mutagenesis, a well-known technique for which protocols and reagents are commercially available (Zoller *et al.*, *Nucleic Acids Res.* **1982**, 10:6487-6500 and Adelman *et al.*, *DNA* **1983**, 2:183-193)). The isolated DNA is analyzed by restriction and/or sequenced by the dideoxy nucleotide method of Sanger, *Proc. Natl. Acad. Sci. USA* **1977**, 74:5463) as further described by Messing, *et al.*, *Nucleic Acids Res.* **1981**, 9:309, or by the method of Maxam *et al.*, *Meth. Enzymol.*, *supra*.

Vector DNA can be introduced into mammalian cells via conventional techniques such as calcium phosphate or calcium chloride co-precipitation, DEAE-dextran-mediated transfection, lipofection, or electroporation. Suitable methods for transforming host cells can be found in Sambrook *et al. supra* and other standard texts. In fusion expression vectors, a proteolytic cleavage site is introduced at the junction of the reporter group and the target protein to enable separation of the target protein from the reporter group subsequent to purification of the fusion protein. Proteolytic enzymes for such cleavage and their recognition sequences include Factor Xa, thrombin and enterokinase.

#### 4.6 Therapeutic Uses

In accordance with the invention, a compound of the invention and/or a pharmaceutical composition thereof is administered to a patient, preferably a human, suffering from cancer or from a disease characterized by angiogenesis. The compounds of the invention and/or pharmaceutical compositions thereof may be used to treat or prevent cancer or undesired angiogenesis.

Preferably, cancers include any vascularized tumor, preferably, a solid tumor, including but not limited to, carcinomas of the lung, breast, ovary, stomach, pancreas, larynx, esophagus, testes, liver, parotid, biliary tract, colon, rectum, cervix, uterus, endometrium, kidney, bladder, prostate, thyroid, squamous cell carcinomas, adenocarcinomas, small cell carcinomas, melanomas, gliomas, neuroblastomas, sarcomas (*e.g.*, angiosarcomas, chondrosarcomas)). Disease characterized by undesired angiogenesis include, but are not limited to, arthritis, diabetes, arteriosclerosis, arteriovenous, malformations, corneal graft neovascularization, delayed wound healing, diabetic retinopathy, age related macular degeneration, granulations, burns, hemophilic joints, rheumatoid arthritis, hypertrophic scars, neovascular glaucoma, nonunion fractures, Osier Weber Syndrome, psoriasis, pyogenic, granuloma, retrolental fibroplasia, pterygium, scleroderma, trachoma, vascular adhesions, ocular neovascularization, parasitic diseases, hypertrophy following surgery, inhibition of hair growth, macular degeneration (including both wet and dry type), rheumatoid arthritis and osteoarthritis.

Also contemplated are methods for treating a patient having a disease or condition associated with undesired cell migration, invasion or proliferation comprising administering to the subject an therapeutically effective amount of a

compound of the invention and/or a pharmaceutical composition thereof. In the above methods, the patient has a tumor, and angiogenesis inhibition results in reduction in size or growth rate of the tumor or destruction of the tumor. Preferably, the subject is a human.

5           Other examples of diseases or conditions against which the above methods may be effective include primary growth of a solid tumor, leukemia or lymphoma, tumor invasion, metastasis or growth of tumor metastases; benign hyperplasia; atherosclerosis, myocardial angiogenesis; post-balloon angioplasty vascular restinosis, neointima formation following vascular trauma, vascular graft restinosis, coronary  
10 collateral formation, deep venous thrombosis, ischemic limb angiogenesis; telangiectasia, pyogenic granuloma, corneal disease, rubeosis, neovascular glaucoma, diabetic and other retinopathy, retrolental fibroplasias, diabetic neovascularization, macular degeneration, endometriosis, arthritis, fibrosis associated with a chronic inflammatory condition, traumatic spinal cord injury including ischemia, scarring or  
15 fibrosis, lung fibrosis, chemotherapy-induced fibrosis; wound healing with scarring and fibrosis, peptic ulcers, a bone fracture, keloids, or a disorder of vasculogenesis, hematopoiesis, ovulation, menstruation, pregnancy or placentation associated with pathogenic cell invasion or with angiogenesis.

          A preferred disease or condition to be treated by the above methods are tumor  
20 growth, invasion or metastasis, particularly brain tumors. Examples of such brain tumors are astrocytoma, anaplastic astrocytoma, glioblastoma, glioblastoma multiformae, pilocytic astrocytoma, pleiomorphic xanthoastrocytoma, subependymal giant cell astrocytoma, fibrillary astrocytoma, gemistocytic astrocytoma, protoplasmic astrocytoma, oligodendroglioma, anaplastic oligodendroglioma, ependymoma,  
25 anaplastic ependymoma, myxopapillary ependymoma, subependymoma, mixed oligoastrocytoma and malignant oligoastrocytoma.

          The above methods may also be used to treat uterine diseases such as endometriosis and pathogenic ocular neovascularization associated with, or a cause of, proliferative diabetic retinopathy, neovascular age-related macular degeneration,  
30 retinopathy of prematurity, sickle cell retinopathy or retinal vein occlusion.

          Further, in certain embodiments, compounds of the invention and/or pharmaceutical compositions thereof are administered to a patient, preferably a human, as a preventative measure against various diseases or disorders characterized by undesired angiogenesis including cancer. Thus, the compounds of the invention

and/or pharmaceutical compositions thereof may be administered as a preventative measure to a patient having a predisposition for a disease characterized by undesired angiogenesis. Accordingly, the compounds and/or pharmaceutical compositions of the invention may be used for the prevention of one disease or disorder and  
5 concurrently treating another (*e.g.*, preventing arthritis while treating cancer).

The suitability of the compounds of the invention and/or pharmaceutical compositions thereof in treating or preventing various diseases or disorders such as cancer may be assayed by methods described herein and in the art. Accordingly, it is well with the capability of those of skill in the art to assay and use the compounds of  
10 the invention and/or pharmaceutical compositions thereof to treat or prevent diseases or disorders such as cancer.

#### 4.7 Diagnostic Uses and Methods

A compound of the invention and/or a pharmaceutical composition thereof is  
15 administered to a patient, preferably a human, in a diagnostically effective amount to detect or image a disease such as those listed in Section 5.6 above. Further, compounds of the invention and/or pharmaceutical compositions thereof may be used to detect or image diseases or conditions associated with undesired cell migration, invasion or proliferation such as those listed above in Section 5.6 by administering to  
20 a subject an diagnostically effective amount of a compound of the invention and/or a pharmaceutical composition thereof.

Compounds of the invention may be diagnostically labeled and used, for example, to detect cell migration, cell invasion and cell proliferation. The disposition of a compound of the invention during and after binding may be followed *in vitro* or  
25 *in vivo* by using an appropriate method to detect the label. Diagnostically labeled compounds may be utilized *in vivo* for diagnosis and prognosis, for example, to image occult metastatic foci or for other types of *in situ* evaluations. For diagnostic applications, compounds of the invention may include bound linker moieties, which are well known to those of skill in the art

30 *In situ* detection of the labeled compound may be accomplished by removing a histological specimen from a subject and examining it by microscopy under appropriate conditions to detect the label. Those of ordinary skill will readily perceive that any of a wide variety of histological methods (such as staining procedures) can be modified in order to achieve such *in situ* detection.

For diagnostic *in vivo* radioimaging, the type of detection instrument available is a major factor in selecting a radionuclide. The radionuclide chosen must have a type of decay which is detectable by a particular instrument. In general, any conventional method for visualizing diagnostic imaging can be utilized in accordance with this invention. Another factor in selecting a radionuclide for *in vivo* diagnosis is that its half-life be long enough so that the label is still detectable at the time of maximum uptake by the target tissue, but short enough so that deleterious irradiation of the host is minimized. In one preferred embodiment, a radionuclide used for *in vivo* imaging does not emit particles, but produces a large number of photons in a 140-200 keV range, which may be readily detected by conventional gamma cameras.

*In vivo* imaging may be used to detect occult metastases which are not observable by other methods. Compounds of the present invention may be used in diagnostic, prognostic or research procedures in conjunction with any appropriate cell, tissue, organ or biological sample of a desired animal species. By the term “biological sample” is intended any fluid or other material derived from the body of a normal or diseased subject, such as blood, serum, plasma, lymph, urine, saliva, tears, cerebrospinal fluid, milk, amniotic fluid, bile, ascites fluid, pus and the like. Also included within the meaning of this term is a organ or tissue extract and a culture fluid in which any cells or tissue preparation from the subject has been incubated.

Useful doses are defined as effective amount of a compound for the particular diagnostic measurement. Thus, an effective amount means an amount sufficient to be detected using the appropriate detection system *e.g.*, magnetic resonance imaging detector, gamma camera, *etc.* The minimum detectable amount will depend on the ratio of labeled compound specifically bound to a tumor (signal) to the amount of labeled compound either bound non-specifically or found free in plasma or in extracellular fluid.

The amount of a composition to be administered depends on the precise compound selected, the disease or condition, the route of administration, and the judgment of the skilled imaging professional. Generally, the amount of a compound needed for detectability in diagnostic use will vary depending on considerations such as age, condition, sex, and extent of disease in the patient, contraindications, if any, and other variables, and is to be adjusted by the individual physician or diagnostician. Dosage can vary from 0.01 mg/kg to 100 mg/kg.



#### 4.8 Therapeutic/Prophylactic Administration

The compounds the invention and/or pharmaceutical compositions thereof may be advantageously used in human medicine. As previously described in Section 4.6 above, compounds of the invention and/or pharmaceutical compositions thereof  
5 are useful for the treatment or prevention of various diseases or disorders such as cancer.

When used to treat or prevent the above disease or disorders, compounds of the invention and/or pharmaceutical compositions thereof may be administered or applied singly, or in combination with other agents. The compounds of the invention  
10 and/or pharmaceutical compositions thereof may also be administered or applied singly, in combination with other pharmaceutically active agents (*e.g.*, other anti-cancer agents, other anti-angiogenic agents such as chelators as zinc, penicillamine, thiomolybdate *etc.*), including other compounds of the invention.

The current invention provides methods of treatment and prophylaxis by  
15 administration to a patient of a therapeutically effective amount of a compound and/or pharmaceutical composition of the invention. The patient may be an animal, is more preferably, a mammal and most preferably, a human.

The present compounds and/or pharmaceutical compositions of the invention, may be administered orally. The compounds and/or pharmaceutical compositions of  
20 the invention may also be administered by any other convenient route, for example, by infusion or bolus injection, by absorption through epithelial or mucocutaneous linings (*e.g.*, oral mucosa, rectal and intestinal mucosa, *etc.*). Administration can be systemic or local. Various delivery systems (*e.g.*, encapsulation in liposomes, microparticles, microcapsules, capsules, *etc.*) may be used to administer a compound  
25 and/or pharmaceutical composition of the invention. Methods of administration include, but are not limited to, intradermal, intramuscular, intraperitoneal, intravenous, subcutaneous, intranasal, epidural, oral, sublingual, intranasal, intracerebral, intravaginal, transdermal, rectally, by inhalation, or topically, particularly to the ears, nose, eyes, or skin. The preferred mode of administration is  
30 left to the discretion of the practitioner, and will depend in-part upon the site of the medical condition. In most instances, administration will result in the release of the compounds and/or pharmaceutical compositions of the invention into the bloodstream.

In specific embodiments, it may be desirable to administer one or more compounds and/or pharmaceutical composition of the invention locally to the area in need of treatment. This may be achieved, for example, and not by way of limitation, by local infusion during surgery, topical application, *e.g.*, in conjunction with a wound dressing after surgery, by injection, by means of a catheter, by means of a suppository, or by means of an implant, said implant being of a porous, non-porous, or gelatinous material, including membranes, such as sialastic membranes, or fibers. In one embodiment, administration can be by direct injection at the site (or former site) of cancer or arthritis.

In certain embodiments, it may be desirable to introduce one or more compounds and/or pharmaceutical compositions of the invention into the central nervous system by any suitable route, including intraventricular, intrathecal and epidural injection. Intraventricular injection may be facilitated by an intraventricular catheter, for example, attached to a reservoir, such as an Ommaya reservoir.

A compound and/or pharmaceutical composition of the invention may also be administered directly to the lung by inhalation. For administration by inhalation, a compound and/or pharmaceutical composition of the invention may be conveniently delivered to the lung by a number of different devices. For example, a Metered Dose Inhaler (“MDI”), which utilizes canisters that contain a suitable low boiling propellant, (*e.g.*, dichlorodifluoromethane, trichlorofluoromethane, dichlorotetrafluoroethane, carbon dioxide or any other suitable gas) may be used to deliver compounds of the invention directly to the lung.

Alternatively, a Dry Powder Inhaler (“DPI”) device may be used to administer a compound and/or pharmaceutical composition of the invention to the lung. DPI devices typically use a mechanism such as a burst of gas to create a cloud of dry powder inside a container, which may then be inhaled by the patient. DPI devices are also well known in the art. A popular variation is the multiple dose DPI (“MDDPI”) system, which allows for the delivery of more than one therapeutic dose. MDDPI devices are available from companies such as AstraZeneca, GlaxoWellcome, IVAX, Schering Plough, SkyePharma and Vectura. For example, capsules and cartridges of gelatin for use in an inhaler or insufflator may be formulated containing a powder mix of a compound of the invention and a suitable powder base such as lactose or starch for these systems.

Another type of device that may be used to deliver a compound and/or pharmaceutical composition of the invention to the lung is a liquid spray device supplied, for example, by Aradigm Corporation, Hayward, CA. Liquid spray systems use extremely small nozzle holes to aerosolize liquid drug formulations that may then  
5 be directly inhaled into the lung.

In one embodiment, a nebulizer is used to deliver a compound and/or pharmaceutical composition of the invention to the lung. Nebulizers create aerosols from liquid drug formulations by using, for example, ultrasonic energy to form fine particles that may be readily inhaled (see *e.g.*, Verschoyle *et al.*, *British J. Cancer*,  
10 1999, 80, Suppl. 2, 96, which is herein incorporated by reference). Examples of nebulizers include devices supplied by Batelle Pulmonary Therapeutics (Columbus, OH) (See, Armer *et al.*, United States Patent No. 5,954,047; van der Linden *et al.*, United States Patent No. 5,950,619; van der Linden *et al.*, United States Patent No. 5,970,974).

In another embodiment, an electrohydrodynamic (“EHD”) aerosol device is used to deliver a compound and/or pharmaceutical composition of the invention to the lung. EHD aerosol devices use electrical energy to aerosolize liquid drug solutions or suspensions (see *e.g.*, Noakes *et al.*, United States Patent No. 4,765,539). The electrochemical properties of the formulation may be important parameters to  
20 optimize when delivering a compound and/or pharmaceutical composition of the invention to the lung with an EHD aerosol device and such optimization is routinely performed by one of skill in the art. EHD aerosol devices may more efficiently deliver drugs to the lung than existing pulmonary delivery technologies.

In another embodiment, the compounds and/or pharmaceutical compositions of the invention can be delivered in a vesicle, in particular a liposome (see Langer,  
25 1990, *Science*, 249:1527-1533; Treat *et al.*, in “Liposomes in the Therapy of Infectious Disease and Cancer,” Lopez-Berestein and Fidler (eds.), Liss, New York, pp.353-365 (1989); see generally “Liposomes in the Therapy of Infectious Disease and Cancer,” Lopez-Berestein and Fidler (eds.), Liss, New York, pp.353-365 (1989)).

In another embodiment, the compounds and/or pharmaceutical compositions of the invention can be delivered *via* sustained release systems, preferably, oral sustained release systems. In one embodiment, a pump may be used (Langer, *supra*; Sefton, 1987, *CRC Crit. Ref. Biomed. Eng.* 14:201; Saudek *et al.*, 1989, *N. Engl. J Med.* 321:574).

In another embodiment, polymeric materials can be used (*see* “Medical Applications of Controlled Release,” Langer and Wise (eds.), CRC Pres., Boca Raton, Florida (1974); “Controlled Drug Bioavailability,” Drug Product Design and Performance, Smolen and Ball (eds.), Wiley, New York (1984); Langer *et al.*, **1983**, *J. Macromol. Sci. Rev. Macromol Chem.* 23:61; Levy *et al.*, **1985**, *Science* 228: 190; During *et al.*, **1989**, *Ann. Neurol.* 25:351; Howard *et al.*, **1989**, *J. Neurosurg.* 71:105).

In another embodiment, polymeric materials are used for oral sustained release delivery. Preferred polymers include sodium carboxymethylcellulose, hydroxypropylcellulose, hydroxypropylmethylcellulose and hydroxyethylcellulose (most preferred, hydroxypropyl methylcellulose). Other preferred cellulose ethers have been described (Alderman, *Int. J. Pharm. Tech. & Prod. Mfr.*, **1984**, 5(3) 1-9). Factors affecting drug release are well known to the skilled artisan and have been described in the art (Bamba *et al.*, *Int. J. Pharm.*, **1979**, 2, 307).

In another embodiment, enteric-coated preparations can be used for oral sustained release administration. Preferred coating materials include polymers with a pH-dependent solubility (*i.e.*, pH-controlled release), polymers with a slow or pH-dependent rate of swelling, dissolution or erosion (*i.e.*, time-controlled release), polymers that are degraded by enzymes (*i.e.*, enzyme-controlled release) and polymers that form firm layers that are destroyed by an increase in pressure (*i.e.*, pressure-controlled release).

In still another embodiment, osmotic delivery systems are used for oral sustained release administration (Verma *et al.*, *Drug Dev. Ind. Pharm.* **2000**, 26:695-708). In still another embodiment, OROS<sup>TM</sup> osmotic devices are used for oral sustained release delivery devices (Theeuwes *et al.*, United States Patent No. 3,845,770; Theeuwes *et al.*, United States Patent No. 3,916,899).

In yet another embodiment, a controlled-release system can be placed in proximity of the target of the compounds and/or pharmaceutical composition of the invention, thus requiring only a fraction of the systemic dose (*see, e.g.*, Goodson, in “Medical Applications of Controlled Release,” *supra*, vol. 2, pp. 115-138 (1984)). Other controlled-release systems discussed in Langer, **1990**, *Science* 249:1527-1533 may also be used.

#### 4.9 Pharmaceutical Compositions

The present pharmaceutical compositions contain a therapeutically or diagnostically effective amount of one or more compounds of the invention, preferably, in purified form, together with a suitable amount of a pharmaceutically acceptable vehicle, so as to provide the form for proper administration to a patient. When administered to a patient, the compounds of the invention and pharmaceutically acceptable vehicles are preferably sterile. Water is a preferred vehicle when the compounds of the invention are administered intravenously. Saline solutions and aqueous dextrose and glycerol solutions can also be employed as liquid vehicles, particularly for injectable solutions. Suitable pharmaceutical vehicles also include excipients such as starch, glucose, lactose, sucrose, gelatin, malt, rice, flour, chalk, silica gel, sodium stearate, glycerol monostearate, talc, sodium chloride, dried skim milk, glycerol, propylene, glycol, water, ethanol and the like. The present pharmaceutical compositions, if desired, can also contain minor amounts of wetting or emulsifying agents, or pH buffering agents. In addition, auxiliary, stabilizing, thickening, lubricating and coloring agents may be used.

Pharmaceutical compositions comprising a compound of the invention may be manufactured by means of conventional mixing, dissolving, granulating, dragee-making, levigating, emulsifying, encapsulating, entrapping or lyophilizing processes. Pharmaceutical compositions may be formulated in conventional manner using one or more physiologically acceptable carriers, diluents, excipients or auxiliaries, which facilitate processing of compounds of the invention into preparations which can be used pharmaceutically. Proper formulation is dependent upon the route of administration chosen.

The present pharmaceutical compositions can take the form of solutions, suspensions, emulsion, tablets, pills, pellets, capsules, capsules containing liquids, powders, sustained-release formulations, suppositories, emulsions, aerosols, sprays, suspensions, or any other form suitable for use. In one embodiment, the pharmaceutically acceptable vehicle is a capsule (see *e.g.*, Grosswald *et al.*, United States Patent No. 5,698,155). Other examples of suitable pharmaceutical vehicles have been described in the art (see Remington's Pharmaceutical Sciences, Philadelphia College of Pharmacy and Science, 19th Edition, 1995).

For topical administration, compounds of the invention may be formulated as solutions, gels, ointments, creams, suspensions, *etc.* as is well-known in the art.

Systemic formulations include those designed for administration by injection, *e.g.*, subcutaneous, intravenous, intramuscular, intrathecal or intraperitoneal injection, as well as those designed for transdermal, transmucosal, oral or pulmonary administration. Systemic formulations may be made in combination with a further  
5 active agent that improves mucociliary clearance of airway mucus or reduces mucous viscosity. These active agents include, but are not limited to, sodium channel blockers, antibiotics, N-acetyl cysteine, homocysteine and phospholipids.

In one embodiment, the compounds of the invention are formulated in accordance with routine procedures as a pharmaceutical composition adapted for  
10 intravenous administration to human beings. Typically, compounds of the invention for intravenous administration are solutions in sterile isotonic aqueous buffer. For injection, a compound of the invention may be formulated in aqueous solutions, preferably, in physiologically compatible buffers such as Hanks' solution, Ringer's solution, or physiological saline buffer. The solution may contain formulatory agents  
15 such as suspending, stabilizing and/or dispersing agents. When necessary, the pharmaceutical compositions may also include a solubilizing agent. Pharmaceutical compositions for intravenous administration may optionally include a local anesthetic such as lignocaine to ease pain at the site of the injection. Generally, the ingredients are supplied either separately or mixed together in unit dosage form, for example, as a  
20 lyophilized powder or water free concentrate in a hermetically sealed container such as an ampoule or sachette indicating the quantity of active agent. When the compound of the invention is administered by infusion, it can be dispensed, for example, with an infusion bottle containing sterile pharmaceutical grade water or saline. When a compound of the invention is administered by injection, an ampoule  
25 of sterile water for injection or saline can be provided so that the ingredients may be mixed prior to administration.

For transmucosal administration, penetrants appropriate to the barrier to be permeated are used in the formulation. Such penetrants are generally known in the art.

30 Pharmaceutical compositions for oral delivery may be in the form of tablets, lozenges, aqueous or oily suspensions, granules, powders, emulsions, capsules, syrups, or elixirs, for example. Orally administered pharmaceutical compositions may contain one or more optionally agents, for example, sweetening agents such as fructose, aspartame or saccharin; flavoring agents such as peppermint, oil of

wintergreen, or cherry coloring agents and preserving agents, to provide a pharmaceutically palatable preparation. Moreover, when in tablet or pill form, the compositions may be coated to delay disintegration and absorption in the gastrointestinal tract, thereby providing a sustained action over an extended period of time. Selectively permeable membranes surrounding an osmotically active driving compound are also suitable for orally administered compounds of the invention. In these later platforms, fluid from the environment surrounding the capsule is imbibed by the driving compound, which swells to displace the agent or agent composition through an aperture. These delivery platforms can provide an essentially zero order delivery profile as opposed to the spiked profiles of immediate release formulations. A time delay material such as glycerol monostearate or glycerol stearate may also be used. Oral compositions can include standard vehicles such as mannitol, lactose, starch, magnesium stearate, sodium saccharine, cellulose, magnesium carbonate, *etc.* Such vehicles are preferably of pharmaceutical grade.

For oral liquid preparations such as, for example, suspensions, elixirs and solutions, suitable carriers, excipients or diluents include water, saline, alkyleneglycols (*e.g.*, propylene glycol), polyalkylene glycols (*e.g.*, polyethylene glycol) oils, alcohols, slightly acidic buffers between pH 4 and pH 6 (*e.g.*, acetate, citrate, ascorbate at between about 5.0 mM to about 50.0 mM), *etc.* Additionally, flavoring agents, preservatives, coloring agents, bile salts, acylcarnitines and the like may be added.

For buccal administration, the pharmaceutical compositions may take the form of tablets, lozenges, *etc.* formulated in conventional manner.

Liquid drug formulations suitable for use with nebulizers and liquid spray devices and EHD aerosol devices will typically include a compound of the invention with a pharmaceutically acceptable vehicle. Preferably, the pharmaceutically acceptable vehicle is a liquid such as alcohol, water, polyethylene glycol or a perfluorocarbon. Optionally, another material may be added to alter the aerosol properties of the solution or suspension of compounds of the invention. Preferably, this material is liquid such as an alcohol, glycol, polyglycol or a fatty acid. Other methods of formulating liquid drug solutions or suspension suitable for use in aerosol devices are known to those of skill in the art (see, *e.g.*, Biesalski, United States Patent No. 5,112,598; Biesalski, United States Patent No. 5,556,611).

A compound of the invention may also be formulated in rectal or vaginal pharmaceutical compositions such as suppositories or retention enemas, *e.g.*, containing conventional suppository bases such as cocoa butter or other glycerides.

In addition to the formulations described previously, a compound of the invention may also be formulated as a depot preparation. Such long acting formulations may be administered by implantation (for example, subcutaneously or intramuscularly) or by intramuscular injection. Thus, for example, a compound of the invention may be formulated with suitable polymeric or hydrophobic materials (for example as an emulsion in an acceptable oil) or ion exchange resins, or as sparingly soluble derivatives, for example, as a sparingly soluble salt.

When a compound of the invention is acidic, it may be included in any of the above-described formulations as the free acid or a pharmaceutically acceptable salt. Pharmaceutically acceptable salts substantially retain the activity of the free acid, may be prepared by reaction with bases and tend to be more soluble in aqueous and other protic solvents than the corresponding free acid form.

#### 4.10 Doses

A compound of the invention, or pharmaceutical compositions thereof, will generally be used in an amount effective to achieve the intended purpose. For use to treat or prevent diseases or disorders such as cancer the compounds of structural Formula (I) and/or pharmaceutical compositions thereof, are administered or applied in a therapeutically effective amount. For use to detect diseases or disorders such as cancer the compounds of structural Formula (I) and/or pharmaceutical compositions thereof, are administered or applied in a diagnostically effective amount.

The amount of a compound of the invention that will be effective in the treatment, prevention or detection of a particular disorder or condition disclosed herein will depend on the nature of the disorder or condition, and can be determined by standard clinical techniques known in the art as previously described. In addition, *in vitro* or *in vivo* assays may optionally be employed to help identify optimal dosage ranges. The amount of a compound of the invention administered will, of course, be dependent on, among other factors, the subject being treated, the weight of the subject, the severity of the affliction, the manner of administration and the judgment of the prescribing physician.



For example, the dosage may be delivered in a pharmaceutical composition by a single administration, by multiple applications or controlled release. In one embodiment, the compounds of the invention are delivered by oral sustained release administration. Preferably, in this embodiment, the compounds of the invention are administered twice per day (more preferably, once per day). Dosing may be repeated intermittently, may be provided alone or in combination with other drugs and may continue as long as required for effective treatment of the disease state or disorder.

Suitable dosage ranges for oral administration are dependent on the potency of the drug, but are generally about 0.001 mg to about 200 mg of a compound of the invention per kilogram body weight. Dosage ranges may be readily determined by methods known to the artisan of ordinary skill.

Suitable dosage ranges for intravenous (i.v.) administration are about 0.01 mg to about 100 mg per kilogram body weight. Suitable dosage ranges for intranasal administration are generally about 0.01 mg/kg body weight to about 1 mg/kg body weight. Suppositories generally contain about 0.01 milligram to about 50 milligrams of a compound of the invention per kilogram body weight and comprise active ingredient in the range of about 0.5% to about 10% by weight. Recommended dosages for intradermal, intramuscular, intraperitoneal, subcutaneous, epidural, sublingual or intracerebral administration are in the range of about 0.001 mg to about 200 mg per kilogram of body weight. Effective doses may be extrapolated from dose-response curves derived from *in vitro* or animal model test systems. Such animal models and systems are well-known in the art.

The compounds of the invention are preferably assayed *in vitro* and *in vivo*, as described above, for the desired therapeutic, prophylactic or diagnostic activity, prior to use in humans. For example, *in vitro* assays can be used to determine whether administration of a specific compound of the invention or a combination of compounds of the invention is preferred for treating, preventing or diagnosing cancer. The compounds of the invention may also be demonstrated to be effective and safe using animal model systems.

Preferably, a therapeutically effective dose of a compound of the invention described herein will provide therapeutic benefit without causing substantial toxicity. Similarly, a diagnostically effective dose of a compound of the invention described herein will provide diagnostic benefit without causing substantial toxicity. Toxicity of compounds of the invention may be determined using standard pharmaceutical

procedures and may be readily ascertained by the skilled artisan. The dose ratio between toxic and therapeutic effect is the therapeutic index. A compound of the invention will preferably exhibit particularly high therapeutic indices in treating disease and disorders. The dosage of a compound of the inventions described herein  
5 will preferably be within a range of circulating concentrations that include an effective therapeutic or diagnostic dose with little or no toxicity.

#### **4.11 Combination Therapy**

In certain embodiments of the present invention, the compounds and/or  
10 pharmaceutical compositions of the invention can be used in combination therapy with at least one other therapeutic agent. The compound and/or pharmaceutical composition of the invention and the therapeutic agent can act additively or, more preferably, synergistically. In one embodiment, a compound and/or pharmaceutical composition of the invention is administered concurrently with the administration of  
15 another therapeutic agent, which may be part of the same pharmaceutical composition or a different pharmaceutical composition. In another embodiment, a pharmaceutical composition of the invention is administered prior or subsequent to administration of another therapeutic agent.

In particular, in one preferred embodiment, the compounds and/or  
20 pharmaceutical compositions of the invention can be used in combination therapy with other chemotherapeutic agents (*e.g.*, alkylating agents (*e.g.*, nitrogen mustards (*e.g.*, cyclophosphamide, ifosfamide, mechlorethamine, melphalen, chlorambucil, hexamethylmelamine, thiotepa), alkyl sulfonates (*e.g.*, busulfan), nitrosoureas, triazines), antimetabolites (*e.g.*, folic acid analogs, pyrimidine analogs (*e.g.*,  
25 fluorouracil, floxuridine, cytosine arabinoside, *etc.*), purine analogs (*e.g.*, mercaptopurine, thioguanine, pentostatin, *etc.*), natural products (*e.g.*, vinblastine, vincristine, etoposide, tertiposide, dactinomycin, daunorubicin, doxorubicin, bleomycin, mithramycin, mitomycin C, L-asparaginase, interferon alpha), platinum coordination complexes (*e.g.*, *cis*-platinum, carboplatin, *etc.*), mitoxantrone,  
30 hydroxyurea, procarbazine, hormones and antagonists (*e.g.*, prednisone, hydroxyprogesterone caproate, medroxyprogesterone acetate, megestrol acetate, diethylstilbestrol, ethinyl estradiol, tamoxifen, testosterone propionate, fluoxymesterone, flutamide, leuprolide, *etc.*), anti-angiogenesis agents or inhibitors (*e.g.*, angiostatin, retinoic acids and paclitaxel, estradiol derivatives,

thiazolopyrimidine derivatives, *etc.*), apoptosis-inducing agents (*e.g.*, antisense nucleotides that block oncogenes which inhibit apoptosis, tumor suppressors, TRAIL, TRAIL polypeptide, Fas-associated factor 1, interleukin-1 $\beta$ -converting enzyme, phosphotyrosine inhibitors, RXR retinoid receptor agonists, carbostyryl derivatives, *etc.*) and chelators (penicillamine, zinc, trientine, *etc.*)).

#### 4.12 Therapeutic Kits.

The current invention provides therapeutic kits comprising the compounds of the invention or pharmaceutical compositions of the invention. The therapeutic kits may also contain other compounds (*e.g.*, chemotherapeutic agents, natural products, hormones or antagonists, anti-angiogenesis agents or inhibitors, apoptosis-inducing agents or chelators) or pharmaceutical compositions of these other compounds.

Therapeutic kits may have a single containers which contains the compound of the invention or pharmaceutical compositions of the invention with or without other components (*e.g.*, other compounds or pharmaceutical compositions of these other compounds) or may have distinct container for each component. Preferably, therapeutic kits of the invention include a compound and/or a pharmaceutical composition of the invention packaged for use in combination with the co-administration of a second compound (preferably, a chemotherapeutic agent, a natural product, a hormone or antagonist, a anti-angiogenesis agent or inhibitor, a apoptosis-inducing agent or a chelator) and/or a pharmaceutical composition thereof. The components of the kit may be pre-complexed or each component may be in a separate distinct container prior to administration to a patient.

The components of the kit may be provided in one or more liquid solutions, preferably, an aqueous solution, more preferably, a sterile aqueous solution. The components of the kit may also be provided as solids, which may be converted into liquids by addition of suitable solvents, which are preferably provided in another distinct container.

The container of a therapeutic kit may be a vial, test tube, flask, bottle, syringe, or any other means of enclosing a solid or liquid. Usually, when there is more than one component, the kit will contain a second vial or other container, which allows for separate dosing. The kit may also contain another container for a pharmaceutically acceptable liquid.

Preferably, a therapeutic kit will contain apparatus (*e.g.*, one or more needles, syringes, eye droppers, pipette, *etc.*), which enables administration of the components of the kit.

5

## 5. Examples

The invention is further defined by reference to the following examples, which describe in detail, preparation of compounds of the invention and methods for assaying for biological activity. It will be apparent to those skilled in the art that many modifications, both to materials and methods, may be practiced without departing from the scope of the invention.

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In the examples below, the following abbreviations have the following meanings. If an abbreviation is not defined, it has its generally accepted meaning.

	AcCN	=	acetonitrile
15	Boc	=	<i>tert</i> -butyloxycarbonyl
	CPM	=	counts per minute
	DMF	=	<i>N,N</i> -dimethylformamide
	DMSO	=	dimethylsulfoxide
	Fmoc	=	9-fluorenylmethyloxycarbonyl
20	g	=	gram
	h	=	hour
	HBTU	=	O-Benzotriazole, N, N, N, N, tetramethyl uronium hexafluoro phosphate
	HBSS	=	Hank's buffered saline solution
25	HOBT	=	N-hydroxybenzotriazole
	HPLC	=	high pressure liquid chromatography
	L	=	liter
	LC/MS	=	liquid chromatography/mass spectroscopy
30	M	=	molar
	min	=	minute
	mL	=	milliliter
	mmol	=	millimoles
	NHS	=	N-hydroxysuccinimide
35	NMM	=	N-methyl morpholine
	TFA	=	trifluoroacetic acid
	TIS	=	triisopropylsilane
	TLC	=	thin layer chromatography
	μL	=	microliter
40	μM	=	micromolar
	v/v	=	volume to volume

### 5.1 Example 1: Standard Resin Bound Amino Acid Coupling

Rink Amide AM resin (Novabiochem) was treated with 20% piperidine in DMF (1 mL per 100 mg of resin) for three minutes with nitrogen agitation or vibration and the reaction mixture was filtered. This step was repeated an additional two times. The resin was washed three times with DMF, three times with methanol and three times with dichloromethane. The desired Fmoc protected, tritylated amino acid (BOC or iVDde was used for lysine and t-Bu used for tyrosine) (3 eq), HBTU (3 eq), and HOBt (3 eq) were dissolved in DMF (1 mL per 100 mg of resin) and added to the above resin, followed by the addition of *N*-Methylmorpholine (NMM) (6 eq) and the mixture was agitated for 1 hour. The reaction mixture was filtered and the resin was washed with DMF. This coupling step was repeated and then the resin was washed with DMF three times, MeOH three times and DCM three times. The resin was treated with 20% piperidine in DMF (3 x 3 minutes each) as described above to remove the Fmoc group on the first amino acid. The next desired Fmoc protected, tritylated amino acid (3 eq), HBTU (3 eq), and HOBt (3 eq) were dissolved in DMF (1 mL per 100 mg of resin) and added to the above resin, followed by the addition of *N*-Methylmorpholine (NMM) (6 eq) and the mixture was agitated for 1 hour. The reaction mixture was filtered and the resin was washed with DMF three times, MeOH three times and DCM three times. Subsequent amino acids were single coupled in a similar manner. For peptides containing an N-terminus acetyl, the following commercially available amino acids, Ac-Pro-OH (3 eq) or Ac-Tyr-OH (3 eq) were used and coupled with HBTU (3 eq), HOBt (3 eq) and NMM (6 eq) for an hour. For all other examples, the capping of the N-terminus was performed by coupling the terminal amine on the resin with the appropriate acid (3 eq), HBTU (3 eq), HOBt (3 eq) and NMM (6 eq). The fully assembled N-capped peptide on Rink Amide AM resin was treated with TFA/TIS/water (95:2.5:2.5, 1 mL per 100 mg of resin) and agitated with nitrogen or vibration for 1 hour. The reaction mixture was filtered, the resin was washed with TFA/TIS/water and with dichloromethane. The solvent was removed *in vacuo* and the resulting residue was triturated three times with ether.

## 30                    **5.2 Example 2: Purification of Peptides**

The crude peptide was dissolved in a minimum amount of methanol and water or in a few drops of glacial acetic acid and water and was purified by preparative reverse phase HPLC (Beckman) with a Phenomenex Synergi hydro-RP C18 column (250mm x 21.2 mm). The peptide was eluted using a gradient from 3-100% B over 30 min

with a flow rate of 20 mL/min, where solvent A was water containing 0.1% trifluoroacetic acid and solvent B was acetonitrile containing 0.1% trifluoroacetic acid. Detection was at 220 or 254 nm. Fractions >95% pure by analytical HPLC analysis (Waters, Phenomenex hydro RP (250mm x 4.6mm over 40 minutes) using  
5 gradient 3-100% B were combined, concentrated to a volume of about 2-4 ml by rotary evaporation, and lyophilized.

### **5.3 Example 3: Cleavage of Peptides from Resin**

The resin was treated with 2% hydrazine in DMF (1 mL per 100 mg of resin)  
10 for three minutes with nitrogen agitation or vibration and the reaction mixture was filtered. This step was repeated an additional five times. The resin was washed three times with DMF and three times with dichloromethane. The desired carboxylic acid (4 eq), HBTU (4 eq) and HOBt (4 eq) were dissolved in DMF (1 mL per 100 mg of resin) and added to the above resin, followed by the addition of *N*-methylmorpholine  
15 (8 eq) and the mixture was agitated for 1 hour. The reaction mixture was filtered and the resin was washed three times with DMF and three times with dichloromethane. The resin was treated with TFA/TIS/water (95:2.5:2.5, 1 mL per 100 mg of resin) and agitated with nitrogen or vibration for 1 hour. The reaction mixture was filtered, the resin was washed once with TFA/TIS/water, three times with dichloromethane and  
20 the filtrate was concentrated. The residue was triturated 4 times with diethyl ether.

### **5.4 Example 4: Coupling of Peptides to Doxorubicin**

25 The desired peptide (1 eq) and doxorubicin hydrochloride (0.6-0.7 eq) were dissolved in DMF (0.05 M) and then HOBt (1.2 eq) and diisopropylethylamine (4 eq) were added to the red solution. The reaction mixture was stirred at room temperature for 5 minutes, and EDAC (1.2 eq) was then added. The reaction mixture was stirred for 17 hours at room temperature and the solvent was removed *in vacuo*.

30

### **5.5 Example 5: 4-Fluorobenzoyl-Pro-His-Ser-Cys-Asn-NH<sub>2</sub>**

This compound was prepared according to the procedure of Examples 1 and 2, with 4-fluorobenzoic acid being used in place of the amino acid. (10.4 mg, 19%) was isolated as a white, fluffy solid: <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>) δ 8.94-8.84 (m, 1H),

8.53-7.96 (m, 4H), 7.66-7.58 (m, 2H), 7.44-7.21 (m, 4H), 7.09 (d,  $J = 10.9$  Hz, 2H), 6.92 (s, 1H), 5.09 (br s, 1H), 4.72-4.65 (m, 1H), 4.52-4.31 (m, 4H), 3.70-3.48 (m, 6H, overlapping with water peak), 3.07-2.72 (m, 4H), 2.22-2.08 (m, 1H), 1.93-1.75 (m, 3H); ES MS  $m/z$  (M+H)<sup>+</sup> 678.4.

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#### **5.6 Example 6: Ac-Pro-His-Ser-Cys(Me)-Asn-OH**

Wang resin (397 mg, 1.00 mmol) was washed three times with dichloromethane, three times with methanol and three times with dichloromethane. Fmoc-Asn(trt)-OH (2.40 g, 4.02 mmol) and diisopropylcarbodiimide (0.625 mL, 4.01 mmol) were dissolved in 4 mL of dichloromethane and 3 mL of DMF at 0°C and stirred for 20 minutes. The dichloromethane was removed *in vacuo*, 3 mL of DMF added to the solution and the solution added to the Wang resin from above in a sintered glass funnel. DMAP (49 mg, 0.40 mmol) in 1 mL DMF was added to the resin and the mixture was agitated with nitrogen for one hour. The mixture was filtered, the resin was washed three times with DMF and three times with dichloromethane, and the coupling procedure was repeated exactly as described above. The remainder of the amino acids were added using the procedure of Example 1, with the exception that the couplings of Fmoc-Ser(trt)-OH and Fmoc-His(trt)-OH were performed using 3 equivalents of amino acid, 3 equivalents of HBTU, 3 equivalents of HOBt, and 6 equivalents of NMM. The procedure of Example 2 was used to cleave and purify the peptide and afforded 282 mg (46%) of a white solid, and as a mixture of two compounds in a ratio of 83:17: <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>)  $\delta$  8.97 (s, 1H), 8.46-8.27 (m, 2H), 8.25-8.15 (m, 1H), 7.98-7.87 (m, 1H), 7.40-7.35 (m, 2H), 6.93 (s, 1H), 4.79-4.60 (m, 1H), 4.56-4.47 (m, 2H), 4.39-4.25 (m, 2H), 3.66-3.60 (m, 3H, overlapping with water peak), 3.22-3.12 (m, 2H), 3.05-2.94 (m, 1H), 2.91-2.82 (m, 1H), 2.68-2.41 (m, 3H, overlapping with DMSO peak), 2.07 (s, 3H), 2.00 (s, 3H), 1.91-1.65 (m, 4H); ES MS  $m/z$  (M+H)<sup>+</sup> 613.4.

#### **5.7 Example 7: Ac-Pro-His-Ser-Cys(Me)-Asn-Gly-Gly-Lys(4-fluorobenzoyl)-NH<sub>2</sub>**

Triethylamine (17  $\mu$ L, 0.12 mmol) was added to a solution of Ac-Pro-His-Ser-Cys(Me)-Asn-Gly-Gly-Lys-NH<sub>2</sub> (21 mg, 0.024 mmol) and 4-fluorobenzoyl succinimide (6.2 mg, 0.026 mmol) in 1 mL of DMF and the solution was stirred at room temperature for 3.5 hours. The reaction mixture was concentrated and the

residue was purified by preparative HPLC (3 to 100% acetonitrile-water over 30 minutes) to give 17 mg (71%) of a white fluffy solid: <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>) δ 8.98 (s, 1H), 8.49-8.42 (m, 1H), 8.35-8.22 (m, 3H), 8.14-8.07 (m, 2H), 7.94-7.87 (m, 3H), 7.82 (d, *J* = 8.0 Hz, 1H), 7.46 (s, 1H), 7.40-7.35 (m, 1H), 7.32-7.26 (m, 3H), 7.02 (s, 2H), 4.82-4.61 (m, 1H), 4.60-4.47 (m, 2H), 4.40-4.26 (m, 2H), 4.21-4.13 (m, 1H), 3.77-3.69 (m, 5H), 3.68-3.61 (m, 2H, overlapping with water peak), 3.27-3.14 (m, 5H, overlapping with water peak), 3.05-2.95 (m, 2H), 2.91-2.83 (m, 1H), 2.72-2.55 (m, 3H, overlapping with DMSO peak), 2.08 (s, 3H), 2.01 (s, 3H), 1.92-1.65 (m, 5H), 1.61-1.46 (m, 3H), 1.38-1.24 (m, 2H); ES MS *m/z* (M+H)<sup>+</sup> 976.7.

10

#### **5.8 Example 8: 5-(& 6-)Carboxyfluorescein-β-Ala-Pro-His-Ser-Cys-Asn-NH<sub>2</sub>**

This compound was prepared from Rink amide AM resin (0.0584 mmol) according to the procedures of Example 1 and 2, with the exceptions that the coupling of Fmoc-Cys(trt)-OH was performed using 3 equivalents of amino acid, 3 equivalents of HBTU, 3 equivalents of HOBT, and 6 equivalents of NMM, while the couplings of Fmoc-Ser(trt)-OH, Fmoc-His(trt)-OH, Fmoc-Pro-OH and 5-(&6-)carboxyfluorescein were performed using 2 equivalents of the acid, 2 equivalents of HBTU, 2 equivalents of HOBT, and 4 equivalents of NMM. The title compound (27 mg, 46%) was isolated as a yellow, fluffy solid, and as a mixture of two compounds: <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>) δ 10.38-10.04 (br s, 1H), 8.98-8.92 (m, 1H), 8.90-8.68 (m, 1H), 8.48-8.41 (m, 1H), 8.35-8.18 (m, 2H), 8.18-8.04 (m, 2H), 8.02-7.96 (m, 1H), 7.66 (s, 0.5H), 7.42-7.33 (m, 2.5H), 7.10 (d, *J* = 7.3 Hz, 2H), 6.95-6.90 (m, 1H), 6.70 (s, 2H), 6.62-6.51 (m, 4H), 4.76-4.57 (m, 1H), 4.51-4.25 (m, 4H), 3.05-2.92 (m, 2H), 2.85-2.71 (m, 3H), 2.69-2.60 (m, 1H), 2.58-2.54 (m, 1H), 2.47-2.37 (m, 2H), 2.09-1.70 (m, 5H); ES MS *m/z* (M+H)<sup>+</sup> 985.6; Anal. calcd for C<sub>45</sub>H<sub>48</sub>N<sub>10</sub>O<sub>14</sub>S: N, 14.22. Found: 10.54 (peptide content, 74%).

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#### **5.9 Example 9: Ac-Pro-His-Ser-Cys-Asn-Gly-Gly-Lys(4-fluorobenzoyl)-NH<sub>2</sub>**

This compound was prepared from Ac-Pro-His(trt)-Ser(trt)-Cys(trt)-Asn(trt)-Gly-Gly-Lys(ivDde)-Rink amide AM resin (208 mg, loading of 0.32 mmol/g, 0.067 mmol) according to Procedure C. The product was cleaved and purified according to Procedure B and afforded 28 mg (44%) of the title compound as a fluffy, white solid: <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>) δ 8.96 (s, 1H), 8.48-8.40 (m, 1H), 8.30-8.18 (m, 2H),

30



8.16-8.00 (m, 2H), 7.97-7.79 (m, 4H), 7.46-7.33 (m, 2H), 7.32-7.24 (m, 4H), 7.11 (s, 0.5H), 7.01 (s, 2H), 6.94 (s, 0.5H), 4.79-4.11 (m, 7H), 3.78-3.60 (m, 8H, overlapping with water peak), 3.27-3.12 (m, 5H, overlapping with water peak), 3.04-2.93 (m, 1H), 2.89-2.70 (m, 2H), 2.64-2.44 (m, 2H, overlapping with DMSO peak), 2.40-2.32 (m, 1H), 2.09-1.97 (m, 3H), 1.91-1.66 (m, 5H), 1.58-1.45 (m, 3H), 1.37-1.23 (m, 2H); ES MS  $m/z$  (M+H)<sup>+</sup> 962.8.

#### **5.10 Example 10: Ac-Tyr-Gly-Gly-Pro-His-Ser-Cys-Asn-NH<sub>2</sub>:**

This compound was prepared from Rink amide AM resin (0.100 mmol) according to the procedures of Examples 1 and 2. The title compound (54 mg, 61%) was isolated as a white, fluffy solid and as a mixture of two compounds in a ratio of 3:1: <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>) δ 8.94 (s, 0.75H), 8.89 (s, 0.25H), 8.54-8.22 (m, 3H), 8.11 (t, *J* = 8.7 Hz, 2H), 7.98 (d, *J* = 7.0 Hz, 1H), 7.84-7.78 (m, 1H), 7.41-7.32 (m, 2H), 7.10 (d, *J* = 8.6 Hz, 2H), 7.05-6.99 (m, 2H), 6.92 (s, 1H), 6.63 (d, *J* = 8.2 Hz, 2H), 4.79-4.55 (m, 13H, overlapping with water peak), 4.49-4.30 (m, 7H, overlapping with water peak), 4.03-3.84 (m, 2H), 3.83-3.58 (m, 4H), 3.55-3.47 (m, 1H), 3.44-3.25 (m, 1H), 3.22-3.09 (m, 1H), 3.06-2.72 (m, 4H), 2.69-2.55 (m, 1H), 2.47-2.38 (m, 2H, overlapping with DMSO peak), 2.09-1.96 (m, 1H), 1.93-1.82 (m, 2H), 1.77 (s, 4H); ES MS  $m/z$  (M+H)<sup>+</sup> 875.7; Anal. calcd for C<sub>36</sub>H<sub>50</sub>N<sub>12</sub>O<sub>12</sub>S: N, 19.21. Found: 12.90 (peptide content, 67%).

#### **5.11 Example 11: Ac-Tyr-Gly-Gly-Gly-Gly-Pro-His-Ser-Cys-Asn-NH<sub>2</sub>**

This compound was prepared from Rink amide AM resin (0.100 mmol) according to the procedures of Examples 1 and 2. The title compound (51 mg, 52%) was isolated as a white, fluffy solid, and as a mixture of two compounds in a ratio of 73:27: <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>) δ 8.94 (s, 0.73H), 8.88 (s, 0.27H), 8.56-8.21 (m, 3H), 8.18-8.01 (m, 4H), 7.99-7.87 (m, 2H), 7.40 (s, 1H), 7.37-7.33 (m, 1H), 7.10 (d, *J* = 8.4 Hz, 2H), 7.02 (d, *J* = 8.5 Hz, 2H), 6.92 (s, 1H), 6.64 (d, *J* = 8.5 Hz, 2H), 4.78-4.26 (m, 26H, overlapping with water peak), 4.03-3.85 (m, 3H, overlapping with water peak), 3.81-3.58 (m, 8H), 3.55-3.46 (m, 1H), 3.21-3.10 (m, 1H), 3.05-2.71 (m, 4H), 2.69-2.56 (m, 1H), 2.51-2.38 (m, 2H, overlapping with DMSO peak), 2.05-1.95

(m, 1H), 1.92-1.82 (m, 2H), 1.77 (s, 4H); ES MS  $m/z$  (M+H)<sup>+</sup> 989.7; Anal. calcd for C<sub>40</sub>H<sub>56</sub>N<sub>14</sub>O<sub>14</sub>S: N, 19.83. Found: 14.52 (peptide content, 73%).

**5.12 Example 12: Ac-Tyr-Gly-Gly-Gly-Gly-Gly-Pro-His-Ser-Cys-Asn-NH<sub>2</sub>**

5 This compound was prepared from Rink amide AM resin (0.100 mmol) according to the procedures of Examples 1 and 2. The title compound (37 mg, 33%) was isolated as a white, fluffy solid, and as a mixture of two compounds in a ratio of 63:37: <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>) δ 8.94 (s, 0.63H), 8.88 (s, 0.37H), 8.55-8.20 (m, 3H), 8.17-7.86 (m, 7H), 7.40 (s, 1H), 7.37-7.32 (m, 1H), 7.10 (d,  $J$  = 8.3 Hz, 2H),  
10 7.02 (d,  $J$  = 8.4 Hz, 2H), 6.92 (s, 1H), 6.63 (d,  $J$  = 8.4 Hz, 2H), 4.76-4.59 (m, 3H, overlapping with water peak), 4.50-4.30 (m, 6H), 4.03-3.85 (m, 2H), 3.81-3.69 (m, 8H), 3.68-3.58 (m, 2H), 3.55-3.47 (m, 2H), 3.43-3.10 (m, 2H), 3.04-2.70 (m, 4H), 2.68-2.55 (m, 2H), 2.47-2.37 (m, 2H, overlapping with DMSO peak), 2.05-1.94 (m, 1H), 1.92-1.81 (m, 2H), 1.77 (s, 4H); ES MS  $m/z$  (M+H)<sup>+</sup> 1103.8; Anal. calcd for  
15 C<sub>44</sub>H<sub>62</sub>N<sub>16</sub>O<sub>16</sub>S: N, 20.32. Found: 14.61 (peptide content, 72%).

**5.13 Example 13: Ac-Tyr-Gly-Gly-Gly-Gly-Gly-Pro-His-Ser-Cys-Asn-NH<sub>2</sub>**

This compound was prepared from Rink amide AM resin (0.100 mmol) according to the procedures of Examples 1 and 2. The title compound (37 mg, 33%)  
20 was isolated as a white, fluffy solid, and as a mixture of two compounds in a ratio of 63:37: <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>) δ 8.94 (s, 0.63H), 8.88 (s, 0.37H), 8.55-8.20 (m, 3H), 8.17-7.86 (m, 7H), 7.40 (s, 1H), 7.37-7.32 (m, 1H), 7.10 (d,  $J$  = 8.3 Hz, 2H), 7.02 (d,  $J$  = 8.4 Hz, 2H), 6.92 (s, 1H), 6.63 (d,  $J$  = 8.4 Hz, 2H), 4.76-4.59 (m, 3H, overlapping with water peak), 4.50-4.30 (m, 6H), 4.03-3.85 (m, 2H), 3.81-3.69 (m,  
25 8H), 3.68-3.58 (m, 2H), 3.55-3.47 (m, 2H), 3.43-3.10 (m, 2H), 3.04-2.70 (m, 4H), 2.68-2.55 (m, 2H), 2.47-2.37 (m, 2H, overlapping with DMSO peak), 2.05-1.94 (m, 1H), 1.92-1.81 (m, 2H), 1.77 (s, 4H); ES MS  $m/z$  (M+H)<sup>+</sup> 1103.8; Anal. calcd for C<sub>44</sub>H<sub>62</sub>N<sub>16</sub>O<sub>16</sub>S: N, 20.32. Found: 14.61 (peptide content, 72%).

30 **5.14 Example 14: Ac-Tyr-NH-(CH<sub>2</sub>)<sub>4</sub>-CO-Pro-His-Ser-Cys-Asn-NH<sub>2</sub>**

This compound was prepared from Rink amide AM resin (0.103 mmol) according to the procedures of Examples 1 and 2. The title compound (53 mg, 60%) was isolated as a white, fluffy solid, and as a mixture of two compounds in a ratio of 66:34: <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>) δ 8.97 (s, 1H), 8.45-8.21 (m, 2H), 8.13 (d,  $J$  =

7.9 Hz, 1H), 8.08-7.96 (m, 2H), 7.93-7.85 (m, 1H), 7.40 (s, 1H), 7.37-7.32 (m, 1H), 7.09 (d,  $J = 10.0$  Hz, 2H), 7.02-6.95 (m, 2H), 6.92 (s, 1H), 6.63 (d,  $J = 8.5$  Hz, 2H), 4.79-4.59 (m, 4H, overlapping with water peak), 4.51-4.27 (m, 7H, overlapping with water peak), 3.72-3.56 (m, 2H), 3.54-3.28 (m, 2H), 3.21-3.10 (m, 1H), 3.07-2.94 (m, 3H), 2.88-2.72 (m, 3H), 2.65-2.55 (m, 1H), 2.46-2.38 (m, 2H, overlapping with DMSO peak), 2.33-2.17 (m, 2H), 2.04-1.95 (m, 1H), 1.93-1.74 (m, 6H), 1.50-1.23 (m, 4H); ES MS  $m/z$  ( $M+H$ )<sup>+</sup> 860.7; Anal. calcd for C<sub>37</sub>H<sub>53</sub>N<sub>11</sub>O<sub>11</sub>S: N, 17.92. Found: 12.40 (peptide content, 69%).

10      **5.15 Example 15: 3-(4-Hydroxyphenyl)propionyl-NH-(CH<sub>2</sub>)<sub>4</sub>-CO-Pro-His-Ser-Cys-Asn-NH<sub>2</sub>**

This compound was prepared from Rink amide AM resin (0.103 mmol) according to the procedures of Examples 1 and 2. The title compound (50 mg, 60%) was isolated as a white, fluffy solid and as a mixture of two compounds in a ratio of 15 7:3: <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>) δ 8.97 (s, 1H), 8.47-8.21 (m, 2H), 8.12 (d,  $J = 7.9$  Hz, 1H), 8.09-7.92 (m, 1H), 7.81-7.73 (m, 1H), 7.40 (s, 1H), 7.37-7.32 (m, 1H), 7.09 (d,  $J = 10.0$  Hz, 2H), 6.99-6.89 (m, 3H), 6.64 (d,  $J = 8.5$  Hz, 2H), 4.78-4.59 (m, 2H), 4.51-4.27 (m, 6H, overlapping with water peak), 3.71-3.56 (m, 3H), 3.54-3.45 (m, 2H), 3.44-3.27 (m, 1H), 3.21-3.10 (m, 1H), 3.06-2.93 (m, 3H), 2.85-2.75 (m, 2H), 20 2.72-2.64 (m, 2H), 2.46-2.37 (m, 2H, overlapping with DMSO peak), 2.32-2.21 (m, 4H), 2.08-1.95 (m, 1H), 1.90-1.69 (m, 3H), 1.52-1.23 (m, 4H); ES MS  $m/z$  ( $M+H$ )<sup>+</sup> 803.6; Anal. calcd for C<sub>35</sub>H<sub>50</sub>N<sub>10</sub>O<sub>10</sub>S: N, 17.45. Found: 12.86 (peptide content, 74%).

**5.16 Example 16: Ac-Pro-His-Ser-Cys(Me)-Asn-Dox**

25      This compound was prepared from Ac-Pro-His-Ser-Cys(Me)-Asn-OH (62 mg, 0.10 mmol) and doxorubicin hydrochloride (37 mg, 0.063 mmol) according to the procedure of Example 4. The resulting residue was purified according to the procedure of Example 2 and afforded the title compound (28 mg, 39%) as a fluffy, orange solid: <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>) δ 8.97-8.93 (m, 1H), 8.44-8.16 (m, 30 3H), 7.96-7.87 (m, 3H), 7.70-7.63 (m, 1H), 7.41-7.26 (m, 3H), 6.93-6.86 (m, 1H), 5.48-5.42 (m, 1H), 5.24-5.20 (m, 1H), 5.10-4.92 (m, 2H), 4.79-4.40 (m, 7H), 4.37-4.24 (m, 2H), 4.16-4.11 (m, 1H), 4.01-3.88 (m, 4H), 3.65-3.48 (m, 6H, overlapping with water peak), 3.25-3.11 (m, 2H), 3.06-2.91 (m, 3H), 2.86-2.70 (m, 1H), 2.67-2.55

(m, 2H, overlapping with DMSO peak), 2.39-2.25 (m, 1H), 2.22-2.07 (m, 2H), 2.04 (s, 3H), 2.00 (s, 3H), 1.90-1.62 (m, 4H), 1.52-1.41 (m, 1H), 1.12 (d,  $J = 6.4$  Hz, 3H); ES MS  $m/z$  (M+H)<sup>+</sup> 1138.5.

5                    **5.17 Example 17: Ac-Pro-His-Ser-Cys(Ac)-Asn-Dox**

This compound was prepared from Ac-Pro-His-Ser-Cys(Ac)-Asn-OH (52 mg, 0.081 mmol) and doxorubicin hydrochloride (33 mg, 0.057 mmol) according to the procedure of Example 3. The resulting residue was purified according to the procedure of Example 2 and afforded the title compound (36 mg, 54%) as a fluffy, orange solid: <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>)  $\delta$  8.95 (s, 1H), 8.44-8.10 (m, 3H), 7.96-7.85 (m, 2H), 7.69-7.62 (m, 1H), 7.40-7.28 (m, 2H), 6.90 (s, 1H), 5.46 (br s, 1H), 5.32-5.21 (m, 1H), 4.98-4.92 (m, 1H), 4.76-4.12 (m, 8H), 4.00-3.88 (m, 3H), 3.07-2.73 (m, 5H), 2.38-2.26 (m, 1H), 2.28 (s, 3H), 2.22-2.06 (m, 2H), 2.00 (s, 3H), 1.91-1.63 (m, 5H), 1.52-1.42 (m, 1H), 1.13 (d,  $J = 6.4$  Hz, 3H); ES MS  $m/z$  (M+H)<sup>+</sup> 1166.6.

**5.18 Example 18: Ac-Pro-His-Ser-Cys(Me)-Asn-Gly-Gly-Lys(CO-(CH<sub>2</sub>)<sub>3</sub>-CO-Dox)-NH<sub>2</sub>**

Ac-Pro-His-Ser-Cys(Me)-Asn-Gly-Gly-Lys-NH<sub>2</sub> (74 mg, 0.087 mmol) and Fmoc-Dox-hemiglutarate (70% pure, 74 mg, 0.059 mmol) were dissolved in 3 mL of DMF at room temperature. HOBt (14 mg, 0.11 mmol) and diisopropylethylamine (60  $\mu$ L, 0.34 mmol) were added to the red solution and the reaction mixture was stirred for 5 minutes. EDAC (21 mg, 0.11 mmol) was added to the reaction mixture and stirred for an additional 18 hours. The solvent was removed *in vacuo* and the resulting red oil was triturated three times with ethyl acetate to afford a red solid. The red solid was dissolved in 5 mL of DMF and 500  $\mu$ L of piperidine. The reaction mixture was cooled to 0°C in an ice bath and a mixture of 450  $\mu$ L TFA, 1.05 mL pyridine and 3mL of DMF was added. After 5 minutes, the solvent was removed *in vacuo* and the residue was triturated once with ethyl acetate and once with diethyl ether to give a red solid. This product was purified according to the procedure of Example 2 and afforded the title compound (14 mg, 16%) as a fluffy, orange solid: ES MS  $m/z$  (M+H)<sup>+</sup> 1493.8.

#### **5.19 Example 19: 2-Fluoroacetyl- $\beta$ -Ala-Pro-His-Ser-Cys-Asn-NH<sub>2</sub>**

This compound was prepared from Rink amide AM resin (0.111 mmol) according to the procedures of Examples 1 and 2. The title compound (30 mg, 40%) was isolated as a white, fluffy solid: <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>)  $\delta$  8.98-8.93 (m, 1H), 8.43-8.21 (m, 2H), 8.15-8.07 (m, 2H), 8.01-7.90 (m, 1H), 7.41-7.32 (m, 2H), 7.09 (d, *J* = 9.9 Hz, 2H), 6.92 (s, 1H), 4.87-4.70 (m, 2H), 4.68-4.59 (m, 1H), 4.50-4.28 (m, 5H), 3.36-3.28 (m, 4H, overlapping with water peak), 3.22-3.10 (m, 2H), 3.05-2.91 (m, 1H), 2.89-2.72 (m, 3H), 2.46-2.36 (m, 3H, overlapping with DMSO peak), 2.09-1.95 (m, 1H), 1.90-1.55 (m, 4H); ES MS *m/z* (*M*+H)<sup>+</sup> 687.5; Anal. calcd for C<sub>26</sub>H<sub>39</sub>FN<sub>10</sub>O<sub>9</sub>S: N, 20.40. Found: 14.89 (peptide content, 73%).

#### **5.20 Example 20: Ac-Pro-His-Ser-Cys-Asn-Gly-Gly-Lys(2-fluoroacetyl)-NH<sub>2</sub>**

This compound was prepared from Ac-Pro-His(trt)-Ser(trt)-Cys(trt)-Asn(trt)-Gly-Gly-Lys(ivDde)-Rink amide AM resin (317 mg, loading of 0.32 mmol/g, 0.101 mmol) according to the procedure of Example 3. The product was purified according to the procedure of Example 2 and afforded 21 mg (23%) of the title compound as a fluffy, white solid: <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>)  $\delta$  8.97 (s, 1H), 8.45-8.17 (m, 3H), 8.16-8.01 (m, 3H), 7.95 (d, *J* = 7.2 Hz, 1H), 7.80 (d, *J* = 8.1 Hz, 1H), 7.45 (s, 1H), 7.41-7.33 (m, 1H), 7.27 (s, 1H), 7.01 (s, 2H), 4.74 (d, *J* = 47 Hz, 2H), 4.78-4.61 (m, 1H), 4.59-4.24 (m, 5H), 4.19-4.10 (m, 2H), 3.55-3.46 (m, 3H, overlapping with water peak), 3.38-2.94 (m, 6H), 2.87-2.70 (m, 2H), 2.65-2.42 (m, 2H, overlapping with DMSO peak), 2.36 (t, *J* = 8.5 Hz, 1H), 2.00 (s, 3H), 1.90-1.61 (m, 5H), 1.56-1.36 (m, 3H), 1.31-1.15 (m, 2H); ES MS *m/z* (*M*+H)<sup>+</sup> 900.8; Anal. calcd for C<sub>35</sub>H<sub>54</sub>FN<sub>13</sub>O<sub>12</sub>S: N, 20.23. Found: 14.55 (peptide content, 72%).

#### **5.21 Example 21: Ac-Pro-His-Ser-Cys-Asn-Gly-Gly-Lys(8-[(4'-fluorobenzyl)amino]suberoyl)-NH<sub>2</sub>**

This compound was prepared from Ac-Pro-His(trt)-Ser(trt)-Cys(trt)-Asn(trt)-Gly-Gly-Lys(ivDde)-Rink amide AM resin (312 mg, loading of 0.32 mmol/g, 0.0997 mmol) according to the procedure of Example 3. The product (42.3 mg of crude material) was purified according to the procedure of Example 2 and afforded 3.9 mg (11%) of PLG-107 as a fluffy, white solid: ES MS *m/z* (*M*+H)<sup>+</sup> 1104.0.

**5.22 Example 22: Ac-Pro-His-Ser-Cys-Asn-β-Ala-Lys(3-(4-hydroxyphenyl)propionyl)-NH<sub>2</sub>**

Ac-Pro-His(trt)-Ser(trt)-Cys(trt)-Asn(trt)-β-Ala-Lys(ivDde) was prepared on Rink amide AM resin (0.100 mmol) according to the procedure of Example 1, but was  
5 not cleaved from the resin. The title compound was prepared from this resin bound peptide according to the procedure of Example 3. The crude product was purified according to the procedure of Example 2 and afforded 39 mg (41%) of the title compound as a fluffy, white solid: <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>) δ 8.97 (s, 1H), 8.46-8.26 (m, 1H), 8.21 (d, *J* = 7.7 Hz, 1H), 8.13 (d, *J* = 7.7 Hz, 1H), 8.07-7.87 (m,  
10 2H), 7.77-7.66 (m, 2H), 7.40 (s, 1H), 7.36-7.28 (m, 2H), 6.99-6.88 (m, 4H), 6.67-6.62 (m, 2H), 4.80-4.60 (m, 1H), 4.52-4.25 (m, 6H), 3.72-3.59 (m, 4H, overlapping with water peak), 3.56-3.44 (m, 2H), 3.38-3.12 (m, 4H), 3.05-2.93 (m, 3H), 2.87-2.75 (m, 2H), 2.71-2.64 (m, 2H), 2.45-2.35 (m, 2H), 2.31-2.23 (m, 4H), 2.00 (s, 3H), 1.91-1.58 (m, 5H), 1.54-1.41 (m, 1H), 1.39-1.17 (m, 4H); ES MS *m/z* (M+H)<sup>+</sup> 945.8; Anal. calcd for C<sub>41</sub>H<sub>60</sub>N<sub>12</sub>O<sub>12</sub>S: N, 17.79. Found: 12.95 (peptide content, 73%).  
15

**5.23 Example 23: Ac-Pro-His-Ser-Cys-Asn-NH-(CH<sub>2</sub>)<sub>4</sub>-CO-Lys(3-(4-hydroxyphenyl)propionyl)-NH<sub>2</sub>**

Ac-Pro-His(trt)-Ser(trt)-Cys(trt)-Asn(trt)-NH-(CH<sub>2</sub>)<sub>4</sub>-CO-Lys(ivDde) was  
20 prepared on Rink amide AM resin (0.100 mmol) according to the procedure of Example 1, but was not cleaved from the resin. The title compound was prepared from this resin bound peptide according to the procedure of Example 3. The crude product was purified according to the procedure of Example 2 and afforded 34 mg (35%) of the title compound as a fluffy, white solid: <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>) δ 8.97 (s, 1H), 8.45-8.19 (m, 2H), 8.13 (d, *J* = 7.8 Hz, 1H), 8.04-7.92 (m, 1H), 7.81-7.71 (m, 2H), 7.66-7.59 (m, 1H), 7.40 (s, 1H), 7.35 (br s, 1H), 7.29 (br s, 1H), 6.96 (d, *J* = 8.4 Hz, 2H), 6.93 (br s, 2H), 6.64 (d, *J* = 8.4 Hz, 2H), 4.81-4.60 (m, 2H), 4.53-4.24 (m, 8H, overlapping with water peak), 3.72-3.60 (m, 3H, overlapping with water peak), 3.56-3.46 (m, 2H), 3.39-3.25 (m, 1H), 3.24-3.11 (m, 1H), 3.07-2.92 (m,  
25 5H), 2.84-2.72 (m, 2H), 2.71-2.63 (m, 2H), 2.46-2.36 (m, 2H, overlapping with DMSO peak), 2.30-2.24 (m, 2H), 2.16-2.07 (m, 2H), 2.00 (s, 3H), 1.91-1.55 (m, 5H), 1.53-1.16 (m, 9H); ES MS *m/z* (M+H)<sup>+</sup> 973.8; Anal. calcd for C<sub>43</sub>H<sub>64</sub>N<sub>12</sub>O<sub>12</sub>S: N, 17.27. Found: 11.42 (peptide content, 66%).  
30

**5.24 Example 24: Ac-Pro-His-Ser-Cys-Asn-NH-(CH<sub>2</sub>)<sub>6</sub>-CO-Lys(3-(4-hydroxyphenyl)propionyl)-NH<sub>2</sub>**

Ac-Pro-His(trt)-Ser(trt)-Cys(trt)-Asn(trt)-NH-(CH<sub>2</sub>)<sub>6</sub>-CO-Lys(ivDde) was prepared on Rink amide AM resin (0.100 mmol) according to the procedure of Example 1, but was not cleaved from the resin. The title compound was prepared from this resin bound peptide according to the procedure of Example 3. The crude product was purified according to the procedure of Example 2 and afforded 29 mg (29%) of the title compound as a fluffy, white solid: <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>) δ 9.12 (s, 1H), 8.91-8.82 (m, 1H), 8.43-8.26 (m, 2H), 8.03-7.97 (m, 1H), 7.94-7.83 (m, 1H), 7.79-7.71 (m, 2H), 7.41-7.20 (m, 3H), 6.96 (d, *J* = 8.5 Hz, 2H), 6.93-6.87 (m, 2H), 6.64 (d, *J* = 8.5 Hz, 2H), 5.09 (br s, 1H), 4.80-4.57 (m, 1H), 4.48-4.39 (m, 1H), 4.36-4.20 (m, 3H), 4.19-4.09 (m, 1H), 3.69-3.53 (m, 4H), 3.04-2.88 (m, 7H), 2.73-2.63 (m, 3H), 2.45-2.33 (m, 3H, overlapping with DMSO peak), 2.29-2.23 (m, 3H), 2.12-2.05 (m, 3H), 1.99 (s, 3H), 1.91-1.56 (m, 5H), 1.51-1.12 (m, 13 H); ES MS *m/z* (M+H)<sup>+</sup> 1002.1; Anal. calcd for C<sub>45</sub>H<sub>68</sub>N<sub>12</sub>O<sub>12</sub>S: N, 16.79. Found: 10.78 (peptide content, 64%).

**5.25 Example 25: Acetyl-Pro-His-Ser-Cys-Asn-Tyr(3-iodo)-NH<sub>2</sub>**

This compound was prepared according to the procedures of Examples 1 and 2 giving the title compound (32.9 mg, 16%) as a fine white powder. The NMR data indicated a mixture of two species in a ratio of about 80:20: <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>) δ 10.06 (s, 1 H), 8.96 (m, 1 H), 8.14-8.43 (m, 3 H), 7.88-8.05 (m, 2 H), 7.50 (d, 1 H, *J* = 1.8 Hz), 7.40 (s, 3 H), 7.09 (bs, 1 H), 6.98-7.03 (m, 2 H), 6.75 (d, 1 H, *J* = 8.4 Hz), 5.06 (bs, 1 H), 4.60-4.74 (m, 1 H), 4.47-4.54 (m, 1 H), 4.37-4.46 (m, 1 H), 4.22-4.36 (m, 3 H), 3.57-3.69 (m, 2 H), 3.47-3.55 (m, 2 H), 3.11-3.21 (m, 1 H), 2.86-3.04 (m, 3 H), 2.59-2.78 (m, 3 H), 2.35-2.43 (m, 1 H), 2.26-2.31 (t, 1 H, *J* = 7.8 Hz), 2.00 (s, 3 H), 1.69-1.94 (m, 4 H); MS *m/z* (C<sub>32</sub>H<sub>43</sub>IN<sub>10</sub>O<sub>10</sub>S+H)<sup>+</sup> 887.4; Anal. calcd for C<sub>32</sub>H<sub>43</sub>IN<sub>10</sub>O<sub>10</sub>S: N, 15.80. Found: N, 12.30 (peptide content: 78%).

**5.26 Example 26: Acetyl-Pro-His-Ser-Cys-Asn-Tyr(3,5-diiodo)-NH<sub>2</sub>**

This compound was prepared according to the procedures of Examples 1 and 2 giving the title compound (25.0 mg, 11%) as a fine white powder. The NMR data indicated a mixture of two species in a ratio of about 80:20: <sup>1</sup>H NMR (300 MHz,

DMSO-d<sub>6</sub>) δ 9.32 (bs, 1 H), 8.94-8.95 (m, 1 H), 8.14-8.43 (m, 3 H), 7.91-8.04 (m, 2 H), 7.58 (s, 2 H), 7.34-7.42 (m, 3 H), 7.12 (s, 1 H), 6.98 (s, 1 H), 5.06 (bs, 1 H) 4.60-4.79 (m 1, H), 4.39-4.53 (m, 2 H), 4.23-4.35 (3 H), 3.58-3.69 (m, 2 H), 3.47-3.54 (m, 2 H), 3.11-3.21 (m, 1 H), 2.56-3.07 (m, 7 H), 2.37-2.44 (m 1 H), 2.29 (t, 1 H, *J* = 9.7 Hz), 2.00 (s, 3 H), 1.68-1.90 (m, 4 H); MS *m/z* (C<sub>32</sub>H<sub>42</sub>I<sub>2</sub>N<sub>10</sub>O<sub>10</sub>S+H)<sup>+</sup> 1013.4; Anal. calcd for C<sub>32</sub>H<sub>42</sub>I<sub>2</sub>N<sub>10</sub>O<sub>10</sub>S: N, 13.83. Found: N, 11.26 (peptide content: 81%).

#### **5.27 Example 27: Acetyl-Pro-His-Ser-Cys(methyl)-Asn-Gly-Gly-Lys-NH<sub>2</sub>**

This compound was prepared according to the procedures of Examples 1 and giving the title compound as (151.2 mg, 44%) as a fine white powder. The NMR data indicated a mixture of two species in a ratio of about 80:20: <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>) δ 8.96-8.97 (m, 1 H), 8.24-8.46 (m, 3 H), 8.09 (app t, 2 H), 7.79-7.95 (m, 2 H), 7.65 (bs, 3 H), 7.45 (bs, 1 H), 7.34-7.39 (m, 1 H), 7.29 (bs, 1 H), 7.05 (bs, 1 H), 7.00 (bs, 1 H), 5.05 (bs, 1 H), 4.60-4.79 (m, 1 H), 4.47-4.58 (2 H), 4.25-4.40 (m, 2 H), 4.12-4.20 (m, 1 H) 3.69-3.79 (m, 4 H), 3.60-3.66 (m, 3 H), 3.12-3.22 (m, 2 H), 2.94-3.04 (m, 1 H), 2.56-2.89 (m, 6 H), 2.07 (s, 3 H), 2.00 (s, 3 H), 1.64-1.88 (m, 4 H), 1.46-1.58 (m, 3 H), 1.23-1.37 (m, 2 H); MS *m/z* (C<sub>34</sub>H<sub>55</sub>N<sub>13</sub>O<sub>11</sub>S+H)<sup>+</sup> 854.7; Anal. calcd for C<sub>34</sub>H<sub>55</sub>N<sub>13</sub>O<sub>11</sub>S: N, 21.32. Found: N, 14.88 (peptide content: 70%).

#### **5.28 Example 28: 3-(4-Hydroxyphenyl)propionyl-Pro-His-Ser-Cys-Asn-NH<sub>2</sub>**

This compound was prepared according to the procedures of Examples 1 and 2 with the following modification: capping of the N-terminus was carried out using N-succinimidyl -3-(4-hydroxyphenyl)propionate (4.0 eq) and TEA (12.0 eq). The title compound (43.0 mg, 38%) was isolated as a fine white powder. The NMR data indicated a mixture of two species in a ratio of about 80:20: <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>) δ 9.12 (bs, 1 H), 8.87-8.95 (m, 1 H), 8.09-8.39 (m, 4 H), 7.97 (app d, 1 H), 7.39 (s, 1 H), 7.33 (bs, 1 H), 6.99-7.09 (m, 4 H), 6.90-6.92 (2 H), 6.61-6.66 (m, 2 H), 5.09 (bs, 1 H), 4.59-4.77 (m, 2 H), 4.27-4.46 (m, 5 H), 3.57-3.70 (m, 3 H), 3.12-3.19 (m, 1 H), 2.97-3.02 (m, 1 H), 2.66-2.80 (m, 4 H), 2.37-2.45 (m, 2 H), 1.63-2.30 (m, 5 H); MS *m/z* (C<sub>30</sub>H<sub>41</sub>N<sub>9</sub>O<sub>9</sub>S+H)<sup>+</sup> 704.4; Anal. calcd for C<sub>30</sub>H<sub>41</sub>N<sub>9</sub>O<sub>9</sub>S: N, 17.91. Found: N, 13.88 (peptide content: 77%).

#### **5.29 Example 29: Acetyl-Tyr(3,5-diiodo)-Pro-His-Ser-Cys-Asn-NH<sub>2</sub>**



This compound was prepared according to the procedures of Examples 1 and 2 giving the title compound (48.8 mg, 29 %) as a fine white powder. The NMR data indicated a mixture of two species in a ratio of about 80:20: <sup>1</sup>H NMR (300 MHz, MeOD) δ 8.79 (m, 1 H), 8.26-8.46 (m, 1 H), 7.61-7.65 (m, 2 H), 7.42-7.47 (m, 1 H), 4.70-4.76 (m, 1 H), 4.52-4.56 (m, 1 H), 4.38-4.45 (m, 2 H), 3.80-3.96 (m, 3 H), 3.59-3.67 (m, 1 H), 3.35-3.43 (m, 2 H), 3.17-3.27 (m, 1 H), 2.99-3.06 (m, 1 H), 2.93-2.95 (m, 2 H), 2.65-2.80 (m, 3 H), 1.99-2.25 (m, 3 H), 1.93-1.96 (m, 3 H); MS *m/z* (C<sub>32</sub>H<sub>42</sub>I<sub>2</sub>N<sub>10</sub>O<sub>10</sub>S+H)<sup>+</sup> 1013.0; Anal. calcd for C<sub>32</sub>H<sub>42</sub>I<sub>2</sub>N<sub>10</sub>O<sub>10</sub>S: N, 13.83. Found: N, 11.33 (peptide content: 82%).

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### **5.30 Example 30: Acetyl-Pro-His-Ser-Cys-Asn-βAla-Tyr(3-iodo)-NH<sub>2</sub>**

This compound was prepared according to the procedures of Examples 1 and 2 giving the title compound (8.1 mg, 6 %) as a fine white powder. The NMR data indicated a mixture of two species in a ratio of about 80:20: <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>) δ 10.07 (s, 1 H), 8.96 (m, 1 H), 8.11-8.44 (m, 3 H), 7.94-8.02 (m, 2 H), 7.68 (app t, 1 H), 7.54 (d, 1 H, *J* = 2.1 Hz), 7.40-7.41 (m, 2 H), 7.34 (m, 1 H), 7.02-7.05 (m, 2 H), 6.90 (bs, 1 H), 6.75 (d, 1 H, *J* = 8.4 Hz), 5.09 (bs, 1 H), 4.60-4.80 (m, 2 H), 4.26-4.51 (m, 4 H), 3.59-3.70 (m, 2 H), 3.47-3.54 (m, 2 H), 3.11-3.21 (m, 4 H), 2.72-3.03 (m, 5 H), 2.56-2.64 (m, 1 H), 2.36-2.43 (m, 2 H), 2.14-2.29 (m, 3 H), 2.00 (s, 3 H), 1.67-1.91 (m, 3 H); MS *m/z* (C<sub>35</sub>H<sub>48</sub>IN<sub>11</sub>O<sub>11</sub>S+H)<sup>+</sup> 958.5; Anal. calcd for C<sub>35</sub>H<sub>48</sub>IN<sub>11</sub>O<sub>11</sub>S: N, 16.09. Found: N, 12.85 (peptide content: 80%).

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### **5.31 Example 31: 3-(4-Hydroxy-3,5-diiodophenyl)propionyl-Pro-His-Ser-Cys-Asn-NH<sub>2</sub>**

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This compound was prepared according to the procedures of Examples 1 and 2 with the following modification: capping of the N-terminus was performed by doing a single coupling with 3-(4-hydroxyphenyl)propionic acid (3.0 eq). The title compound (23.9 mg, 23 %) was isolated as a fine white powder. The NMR data indicated a mixture of two species in a ratio of about 80:20: <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>) δ 9.31 (bs, 1 H), 8.93-8.97 (m, 1 H), 7.99-8.44 (m, 4 H), 7.62 (m, 2 H), 7.41-7.56 (m, 1 H), 7.34 (bs, 1 H), 7.09 (app d, 2 H), 6.92 (s, 1 H), 5.12 (bs, 1 H), 4.29-4.78 (m, 6 H), 3.59-3.72 (m, 3 H), 3.14-3.22 (m, 2 H), 2.96-3.03 (m, 2 H),

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2.77-2.84 (m, 2 H), 2.65-2.71 (m, 2 H), 2.39-2.46 (m, 2 H), 1.56-2.15 (m, 5 H); MS  $m/z$  ( $C_{30}H_{39}I_2N_9O_9S+H$ )<sup>+</sup> 956.3; Anal. calcd for  $C_{30}H_{39}I_2N_9O_9S$ : N, 13.19. Found: N, 10.84 (peptide content: 82%).

5      **5.32 Example 32: Acetyl-Tyr(3,5-diiodo)-β-Ala-Pro-His-Ser-Cys-Asn-NH<sub>2</sub>**

This compound was prepared according to the procedures of Examples 1 and 2 with the following modification: only single couplings were performed with Fmoc-β-Ala-OH and Acetyl-Tyr(3,5-diiodo)-OH. The title compound was isolated as a fine white powder (55.0 mg, 51 %). The NMR data indicated a mixture of two species in a ratio of about 80:20: <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>) δ 9.35 (bs, 1 H), 8.96 (m, 1 H), 8.22-8.41 (m, 3 H), 7.97-8.14 (m, 4 H), 7.60-7.63 (m, 2 H), 7.39 (s, 1 H), 7.34 (bs, 1 H), 7.09 (app d, 2 H), 6.92 (s, 1 H), 4.60-4.78 (m, 2 H), 4.28-4.50 (m, 6 H), 3.59-3.71 (m, 2 H), 2.94-3.06 (m, 2 H), 2.72-2.83 (m, 3 H), 2.38-2.46 (m, 3 H), 1.76-2.09 (m, 7 H); MS  $m/z$  ( $C_{35}H_{47}I_2N_{11}O_{11}S+H$ )<sup>+</sup> 1084.5; Anal. calcd for  $C_{35}H_{47}I_2N_{11}O_{11}S$ : N, 14.22. Found: N, 11.39 (peptide content: 80%).

**5.33 Example 33: Acetyl-Pro-His-Ser-Cys-Asn-β-Ala-Tyr(3,5-diiodo)-NH<sub>2</sub>**

This compound was prepared according to the procedures of Examples 1 and 2 with the following modification: only single couplings were performed with every amino acid. The title compound was isolated as a fine white powder (14.5 mg, 9 %). The NMR data indicated a mixture of two species in a ratio of about 80:20: <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>) δ 9.35 (bs, 1 H), 8.96 (m, 1 H), 7.94-8.44 (m, 6 H), 7.68-7.72 (m, 1 H), 7.62 (s, 2 H), 7.46 (bs, 1 H), 7.40 (bs, 1 H), 7.34 (bs, 1 H), 7.06 (bs, 1 H), 6.90 (bs, 1 H), 5.11 (bs, 1 H), 4.61-4.80 (m, 6 H), 3.57-3.71 (m, 2 H), 3.08-3.22 (m, 3 H), 2.94-3.07 (m, 2 H), 2.73-2.86 (m, 4 H), 2.36-2.43 (m, 2 H), 2.14-2.30 (m, 3 H), 2.00 (s, 3 H), 1.68-1.89 (m, 4 H); MS  $m/z$  ( $C_{35}H_{47}I_2N_{11}O_{11}S$ )<sup>+</sup> 1084.5; Anal. calcd for  $C_{35}H_{47}I_2N_{11}O_{11}S$ : N, 14.22. Found: N, 11.64 (peptide content: 82%).

**5.34 Example 34: Acetyl-Tyr(3-iodo)-Pro-His-Ser-Cys-Asn-NH<sub>2</sub>**

30      This compound was prepared according to the procedures of Example 2 with the following modifications: the final amino acid coupling was carried out with Ac-Tyr(3-iodo)-OH (4 eq), HBTU (4 eq), HOBT (4 eq) and NMM (8 eq). The title compound was isolated (7.7 mg, 16%) as a fine white powder. The NMR data

indicated a mixture of two species in a ratio of about 75:25: <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>) δ 10.10-10.17 (m, 1 H), 8.91-8.97 (m, 1 H), 8.25-8.52-m (2 H), 8.05-8.15 (m, 2 H), 7.58 (d, 1 H, *J* = 2.0 Hz), 7.35-7.43 (m, 2 H), 7.09-7.11 (m, 3 H), 6.92 (bs, 1 H), 6.75-6.77 (m, 1 H), 5.11 (bs, 1 H), 4.55-4.69 (m, 2 H), 4.33-4.50 (m, 4 H), 3.49-  
 5 3.77 (m, 4 H), 3.17 (s, 2 H), 2.69-2.85 (m, 4 H), 1.76-2.07 (m, 6 H) ; MS *m/z* (C<sub>32</sub>H<sub>43</sub>IN<sub>10</sub>O<sub>10</sub>S+H)<sup>+</sup> 887.5; Anal. calcd for C<sub>32</sub>H<sub>43</sub>IN<sub>10</sub>O<sub>10</sub>S: N, 15.80. Found: N, 12.15 (peptide content: 77%).

10 **5.35 Example 35: Acetyl-Pro-His-Ser-Cys-Asn-Gly-Gly-Lys(3-(4-hydroxy-3,4-diiodophenyl) propionyl)-NH<sub>2</sub>**

This compound was prepared according to the procedures of Examples 3 and 2 with the following modifications: the amide formation on lysine was carried out with 3-(4-hydroxy-3,4-diiodophenyl)propionic acid (4 eq) , HBTU (4 eq), HOBT (4 eq) and NMM (8 eq). The title compound was isolated (16.2 mg, 13%) as a fine white  
 15 powder. The NMR data indicated a mixture of two species in a ratio of about 75:25: <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>) δ 9.31 (bs, 1 H), 8.96 (m, 1 H), 8.19-8.44 (m, 3 H), 8.01-8.14 (m, 2 H), 7.95 (m, 1 H), 7.76-7.82 (m, 2 H), 7.55 (s, 2 H), 7.45 (s, 1 H) 7.34, 7.39 (s, 1 H), 7.27 (s, 1 H), 7.01 (s, 2 H), 5.09 (bs, 1 H), 4.61-4.77 (m, 1 H), 4.51-4.57 (m, 1 H), 4.26-4.37 (m, 2 H), 4.10-4.17 (m, 1 H), 3.62-3.74 (m, 5 H), 3.47-  
 20 3.52 (m, 2 H), 2.94-3.02 (m, 5 H), 2.73-2.83 (m, 3 H), 2.64-2.69 (m, 3 H), 2.26-2.39 (m, 4 H), 2.00 (s, 3 H), 1.61-1.89 (m, 4 H), 1.44-1.56 (m, 1 H), 1.18-1.39 (m, 5 H) ; MS *m/z* (C<sub>42</sub>H<sub>59</sub>I<sub>2</sub>N<sub>13</sub>O<sub>13</sub>S+H)<sup>+</sup> 1240.7; Anal. calcd for C<sub>42</sub>H<sub>59</sub>I<sub>2</sub>N<sub>13</sub>O<sub>13</sub>S: N, 14.69. Found: N, 12.28 (peptide content: 84%).

25 **5.36 Example 36: 3-(4-Hydroxy-3,5-diiodophenyl)propionyl-β-Ala-Pro-His-Ser-Cys-Asn-NH<sub>2</sub>**

This compound was prepared according to the procedures of Examples 1 and 2 with the following modification: capping of the N-terminus was carried out using 3-(4-hydroxy-3,5-diiodophenyl)propionic acid (4.0 eq), HBTU (4.0 eq), HOBT (4.0 eq)  
 30 and NMM (8.0 eq). The title compound (23.3 mg, 32 %) was isolated as a fine white powder. The NMR data indicated a mixture of two species in a ratio of about 80:20: <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>) δ 9.32 (bs, 1 H), 8.95 (s, 1 H), 8.22-8.42 (m, 2 H), 8.11-8.13 app d, 1 H), 7.97-7.99 (app d, 1 H), 7.81-7.87 (m, 1 H), 7.56 (s, 2 H), 7.35-

7.39 (m, 2 H), 7.08-7.11 (app d, 2 H), 6.92 (s, 1 H), 5.12 (bs, 1 H), 4.60-4.78 m(m, 1 H), 4.27-4.50 (m, 5 H), 3.59-3.70 (m, 3 H), 2.94-3.02 (m, 1 H), 2.76-2.83 (m, 2 H), 2.65-2.69 (m, 2 H), 2.28-2.46 (m, 6 H), 1.72- 2.09 (m, 4 H); MS  $m/z$  ( $C_{33}H_{44}I_2N_{10}O_{10}S+H$ )<sup>+</sup> 1027.5; Anal. calcd for  $C_{33}H_{44}I_2N_{10}O_{10}S$ : N, 13.64. Found: N, 10.85 (peptide content: 80%).

**5.37 Example 37: Acetyl-Pro-His-Ser-Cys-Asn-Gly-Gly-Lys(5- and 6-carboxyfluorescein)-NH<sub>2</sub>**

This compound was prepared according to the procedures of Examples 2 and 3 with the following modifications: the amide formation on lysine was carried out with 5-(and 6)-carboxyfluorescein (3 eq), HBTU (3 eq), HOBT (3 eq) and NMM (6 eq). The title compound was isolated (10.5 mg, 18%) as a fine white powder. The NMR data indicated a mixture of isomers: <sup>1</sup>H NMR (300 MHz, DMSO- $d_6$ )  $\delta$  10.16 (s, 1 H), 8.78-8.92 (m, 1 H), 8.44, 8.68 (m, m, 1 H), 8.05-8.28 (m, 3 H), 7.79-7.96 (m, 1 H), 7.45, 7.66 (s, s, 1 H), 7.25-7.37 (m, 2 H), 6.97-7.02 (m, 2 H), 6.69 (m, 1 H), 6.52-6.59 (m, 2 H), 5.08 (bs, 1 H), 4.06-4.80 (m, 8 H), 2.24-2.31, 1.99 (m, s, 3 H), 1.67-1.92 (m, 2 H), 1.23-1.59 (m, 3 H); MS  $m/z$  ( $C_{54}H_{63}N_{13}O_{17}S+H$ )<sup>+</sup> 1198.7; Anal. calcd for  $C_{54}H_{63}N_{13}O_{17}S$ : N, 15.20. Found: N, 11.18 (peptide content: 73%).

**5.38 Example 38: (3-(4-Hydroxyphenyl)propionyl)-Gly-Gly-Pro-His-Ser-Cys-Asn-NH<sub>2</sub>**

This compound was prepared according to the procedures of Examples 1 and 2 with the following modification: capping of the N-terminus was carried out using 3-(4-hydroxy-3,5-diiodophenyl)propionic acid (3 eq), HBTU (3 eq), HOBT (3 eq) and NMM (6 eq). The title compound was isolated as a fine white powder (24.2 mg, 30 %). The NMR data indicated a mixture of two species in a ratio of about 80:20: <sup>1</sup>H NMR (300 MHz, DMSO- $d_6$ )  $\delta$  9.17, 8.51 (bs, d, 1 H,  $J$  = 8.2 Hz), 8.87-8.93 (m, 1 H), 8.10-8.31 (m, 3 H), 7.80-7.98 (m, 2 H), 7.35-7.38 (m, 2 H), 7.09 (d, 2 H,  $J$  = 7.4 Hz), 6.96-6.99 (m, 2 H), 6.92 (s, 1 H), 6.63-6.66 (m, 2 H), 5.14 (bs, 1 H), 4.59-4.77 (m, 1 H), 4.30-4.50 (m, 4 H), 3.85-4.00 (m, 2 H), 3.59-3.75 (m, 5 H), 2.89-3.05 (m, 1 H), 2.66-2.83 (m, 4 H), 2.36-2.46 (m, 4 H), 1.97-2.08 (m, 1 H), 1.67-1.90 (m, 2 H); MS  $m/z$  ( $C_{34}H_{47}N_{11}O_{11}S+H$ )<sup>+</sup> 818.6; Anal. Calcd for  $C_{34}H_{47}N_{11}O_{11}S$ : N, 18.84. Found: N, 14.36 (peptide content: 76%).

**5.39 Example 39: Acetyl-Pro-His-Ser-Cys-Asn-Gly-Gly-Lys(3-(4-hydroxyphenyl)propionyl)-NH<sub>2</sub>**

This compound was prepared according to the procedures of Examples 2 and 3 with the following modifications: the amide formation on lysine was carried out with  
5 3-(4-hydroxyphenyl)propionic acid (4 eq), HBTU (4 eq), HOBT (4 eq) and NMM (8 eq). The title compound was isolated (16.2 mg, 13%) as a fine white powder. The NMR data indicated a mixture of two species in a ratio of about 75:25: <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>) δ 8.97 (m, 1 H), 8.42-8.44, 8.19-8.30 (m, m, 3 H), 8.07-8.15 (m, 2 H), 7.94-8.04 (m, 1 H), 7.73-7.82 (m, 2 H), 7.45 (bs, 1 H), 7.35, 7.40 (bs, 1 H), 7.27  
10 (m, 1 H), 6.95-7.01 (m, 4 H), 6.63-6.65 (m, 2 H), 4.72-4.79 (m, 1 H), 4.61-4.68 (m, 1 H), 4.51-4.58 (m, 1 H), 4.41-4.48 (m, 1 H), 4.26-4.39 (m, 2 H), 4.10-4.17 (m, 1 H), 3.61-3.79 (m, 6 H), 3.48-3.55 (m, 2 H), 3.12-3.21 (m, 1 H), 2.96-3.02 (m, 3 H), 2.61-2.86 (m, 4 H), 2.25-2.39 (m, 3 H), 2.00 (s, 3 H), 1.63-1.91 (m, 3 H), 1.44-1.56 (m, 1 H), 1.18-1.37 (m, 4 H); MS *m/z* (C<sub>42</sub>H<sub>61</sub>N<sub>13</sub>O<sub>13</sub>S+H)<sup>+</sup> 988.8; Anal. calcd for  
15 C<sub>42</sub>H<sub>61</sub>N<sub>13</sub>O<sub>13</sub>S: N, 18.43. Found: N, 12.75 (peptide content: 69%).

**5.40 Example 40: (3-(4-Hydroxyphenyl)propionyl)-Gly-Gly-Gly-Gly-Pro-His-Ser-Cys-Asn-NH<sub>2</sub>**

This compound was prepared according to the procedures of Examples 2 and 3  
20 with the following modification: capping of the N-terminus was carried out using 3-(4-hydroxy-3,5-diiodophenyl)propionic acid (3 eq.), HBTU (3 eq.), HOBT (3 eq.) and NMM (6 eq.). The title compound was isolated as a fine white powder (27.3 mg, 29%). The NMR data indicated a mixture of two species in a ratio of about 80:20: <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>) δ 8.87-8.92 (m, 1 H), 8.24, 8.53 (d, d, 1 H, *J* = 7.4, 8.4  
25 Hz), 8.31 (1 H, *J* = 8.4 Hz), 8.08-8.19 (m, 4 H), 7.87-7.98 (m, 2 H), 7.39 (s, 1 H), 7.36 (s, 1 H), 7.10 (d, 1 H, *J* = 6.5 Hz), 6.99 (app d, 2 H), 6.92 (s, 1 H), 6.65 (app d, 2 H), 4.59-4.77 (m, 1 H), 4.30-4.50 (m, 4 H), 3.86-4.02 (m, 2 H), 3.63-3.79 (m, 7 H), 2.92-3.04 (m, 3 H), 2.67-2.84 (m, 6 H), 2.36-2.46 (m, 5 H), 1.97-2.05 (m, 1 H), 1.82-1.90 (m, 2 H), 1.69-1.79 (m, 2 H); MS *m/z* (C<sub>38</sub>H<sub>53</sub>N<sub>13</sub>O<sub>13</sub>S+H)<sup>+</sup> 932.6; Anal. Calcd  
30 for C<sub>38</sub>H<sub>53</sub>N<sub>13</sub>O<sub>13</sub>S: N, 19.54. Found: N, 12.89 (peptide content: 66%).

**5.41 Example 41: (3-(4-Hydroxyphenyl)propionyl)-Gly-Gly-Gly-Gly-Gly-Gly-Pro-His-Ser-Cys-Asn-NH<sub>2</sub>**

This compound was prepared according to the procedures of Examples 1 and 2 with the following modification: capping of the N-terminus was carried out using 3-(4-hydroxy-3,5-diiodophenyl)propionic acid (3 eq), HBTU (3 eq), HOBT (3 eq) and NMM (6 eq). The title compound was isolated as a fine white powder (17.1 mg, 16%). The NMR data indicated a mixture of two species in a ratio of about 70:30: <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>) δ 9.11 (s, 1 H), 8.86, 8.92 (s, s, 1 H), 8.51, 8.05-8.16 (m, d, 6 H, *J* = 8.5 Hz), 8.22-8.32 (m, 2 H) 7.87- 7.98 (m, 2 H), 7.34-7.39 (m, 2 H), 7.11 (s, 1 H), 7.08 (s, 1 H), 6.98 (app d, 2 H), 6.92 (s, 1 H), 6.64 (app d, 2 H), 5.11 (bs, 1 H), 4.59-4.77 (m, 1 H), 4.31-4.50 (m, 4 H), 3.83- 4.02 (m, 2 H), 3.71-3.79 (m, 9 H), 3.58- 3.67 (m, 2 H), 3.48-3.53 (m, 2 H), 3.11-3.19 (m, 1 H), 2.91-3.04 (m, 2 H), 2.76-2.83 (m, 2 H), 2.67-2.72 (m, 2 H), 2.35-2.46 (m, 4 H), 1.96-2.09 (m, 1 H), 1.71-1.90 (m, 4 H); MS *m/z* (C<sub>42</sub>H<sub>59</sub>N<sub>15</sub>O<sub>15</sub>S +H)<sup>+</sup> 1046.8; Anal. Calcd for C<sub>42</sub>H<sub>59</sub>N<sub>15</sub>O<sub>15</sub>S: N, 20.08. Found: N, 15.84 (peptide content: 79%).

**5.42 Example 42: (3-(4-Hydroxyphenyl)propionyl)-Ahp-Pro-His-Ser-Cys-Asn-NH<sub>2</sub>**

This compound was prepared according to the procedures of Examples 1 and 2 with the following modification: capping of the N-terminus was carried out using 3-(4-hydroxyphenyl)propionic acid (3 eq), HBTU (3 eq), HOBT (3 eq) and NMM (6 eq). The title compound was isolated as a fine white powder (47.4 mg, 57%). The NMR data indicated a mixture of two species in a ratio of about 70:30: <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>) δ 8.98 (m, 1 H), 8.21-8.44 (m, 2 H), 7.98-8.14 (m, 2 H), 7.71-7.77 (m, 1 H), 7.40 (s, 1 H), 7.35 (m, 1 H), 7.09 (d, 2 H, *J* = 9.8 Hz), 6.95-6.98 (m, 2 H), 6.92 (s, 1 H), 6.62-6.54 (m, 2 H), 4.62-4.79 (m, 4 H), 4.28-4.50 (m, 8 H), 3.47-3.70 (m, 4 H), 3.12-3.19 (m, 1 H), 2.90-3.03 (m, 3 H), 2.76-2.83 (m, 2 H), 2.65-2.70 (m, 2 H), 2.40-2.46 (m, 2 H), 2.25-2.30 (m, 3 H), 1.72-2.08 (m, 4 H), 1.14-1.50 (m, 7 H); MS *m/z* (C<sub>37</sub>H<sub>54</sub>N<sub>10</sub>O<sub>10</sub>S +H)<sup>+</sup> 831.8; Anal. Calcd for C<sub>37</sub>H<sub>54</sub>N<sub>10</sub>O<sub>10</sub>S: N, 16.86. Found: N, 13.10 (peptide content: 78%).

**5.43 Example 43: Ac-Tyr-Ahp-Pro-His-Ser-Cys-Asn-NH<sub>2</sub>**

This compound was prepared according to the procedures of Examples 1 and 2 with the following modifications: the final amino acid coupling was carried out with Ac-Tyr(3-iodo)-OH (3 eq) , HBTU (3 eq), HOBT (3 eq) and NMM (6 eq). The title compound was isolated (40.9 mg, 46%) as a fine white powder. The NMR data

indicated a mixture of two species in a ratio of about 80:20: <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>) δ 8.97 (m, 1 H), 8.21-8.44 (m, 2 H), 8.06-8.14 (m, 1 H), 7.97-7.99 (m, 2 H), 7.85-7.87 (m, 1 H), 7.40 (s, 1 H), 7.35 (s, 1 H), 7.09 (d, 2 H, J = 9.7 Hz), 6.98-7.00 (m, 2 H), 6.92 (s, 1 H), 6.61-6.64 (m, 2 H), 4.59-4.66 (m, 2 H), 4.25-4.50 (m, 6 H), 3.59-3.71 (m, 2 H), 3.48-3.55 (m, 2 H), 3.12-3.19 (m, 1 H), 2.95-3.10 (m, 3 H), 2.76-2.83 (m, 3 H), 2.57-2.65 (m, 1 H), 2.38-2.46 (m, 2 H), 2.24-2.27 (m, 2 H), 1.80-2.10 (m, 1 H), 1.84-1.89 (m, 2 H), 1.76 (s, 3 H), 1.45-1.49 (m, 2 H), 1.15-1.35 (m, 6 H); MS *m/z* (C<sub>39</sub>H<sub>57</sub>N<sub>11</sub>O<sub>11</sub>S + H)<sup>+</sup> 888.8; Anal. Calcd for C<sub>39</sub>H<sub>57</sub>N<sub>11</sub>O<sub>11</sub>S: N, 17.35. Found: N, 13.81 (peptide content: 80%).

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#### **5.44 Example 44: DOTA-In-β-Ala-Pro-His-Ser-Cys-Asn-NH<sub>2</sub>**

This compound was prepared according to the procedures of Examples 1 and 2 with the following modifications: the final amide coupling was carried out with DOTA-tris(*t*-butyl ester) (3 eq), HBTU (3 eq), HOBT (3 eq) and NMM (6 eq). To the purified peptide-DOTA compound (22.0 mg, 0.022 mmol) in 0.1 M AcOH (aq.) (0.5 mL) was added Indium chloride (48.6 mg, 0.22 mmol) dissolved in 1.0 mL of 0.2 M HCl (aq.) and stirred at rt for 2 hours. The complex was purified according to the procedure of Example 2. The title compound was isolated (40.9 mg, 46%) as a fine white powder. The NMR data indicated a mixture of three species in a ratio of about 70:20:10 <sup>1</sup>H NMR (300 MHz, D<sub>2</sub>O) δ 8.69-8.90 (m, 1 H), 7.39-7.55 (m, 1 H), 4.53-4.64 (m, 3 H), 4.39-4.48 (m, 1 H), 3.88-3.96 (m, 5 H), 3.24-3.68 (m, 28 H), 3.00-3.02 (m, 3 H), 2.69-2.94 (m, 9 H), 2.24-2.43 (m, 2 H), 1.84-2.10 (m, 5 H); MS *m/z* (C<sub>40</sub>H<sub>61</sub>InN<sub>14</sub>O<sub>15</sub>S + H)<sup>+</sup> 11245.8; Anal. Calcd for C<sub>40</sub>H<sub>61</sub>InN<sub>14</sub>O<sub>15</sub>S: N, 17.43. Found: N, 12.42 (peptide content: 71%).

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#### **5.45 Example 45: Acetyl-Pro-His-Ser-Cys-Asn-Gly-Gly-Gly-Gly-Gly-Lys-NH<sub>2</sub>**

This compound was prepared according to the procedures of Examples 1 and 2. The title compound was isolated (15.4 mg, 10%) as a fine white powder. The NMR data indicated a mixture of two species in a ratio of about 75:25: <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>) δ 8.95-8.97 (m, 1 H), 8.43 and 8.02 (d, 1 H, J = 7.6 Hz), 8.11-8.31 (m, 9 H), 7.84-7.86 (m, 1 H), 7.62 (bs, 3 H), 7.44 (bs 1 H), 7.34-7.39 (m, 1 H), 7.29 (s, 1 H), 7.05 (s, 1 H), 7.00 (s, 1 H), 4.51-4.79 (m, 3 H), 4.25-4.48 (m, 4 H), 4.12-4.19

(m, 2 H), 3.74 (m, 10 H), 3.11-3.21 (m, 2 H), 2.94-3.03 (m, 2 H), 2.73-2.82 (m, 5 H), 2.34-2.39 (m, 1 H), 2.00 (s, 3 H), 1.67-1.89 (m, 5 H), 1.46-1.57 (m, 3 H), 1.24-1.34 (m, 2 H); MS  $m/z$  ( $C_{41}H_{65}N_{17}O_{15}S+H$ )<sup>+</sup> 1068.7; Anal. calcd for  $C_{41}H_{65}N_{17}O_{15}S$ : N, 22.29. Found: N, 15.33 (peptide content: 69%).

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**5.46 Example 46: Acetyl-Pro-His-Ser-Cys-Asn-Ava-Tyr-NH<sub>2</sub>**

This compound was prepared according to the procedures of Examples 1 and 2. The title compound was isolated (56.5 mg, 44%) as a fine white powder. The NMR data indicated a mixture of two species in a ratio of about 75:25: <sup>1</sup>H NMR (300  
10 MHz, DMSO-*d*<sub>6</sub>) δ 9.13 (bs, 1 H), 8.96-8.97 (m, 1 H), 8.20-8.30 (m, 2 H), 8.13 (d, 1 H, *J* = 7.8), 7.96-8.04 (m, 1 H), 7.82 (d, 1 H, *J* = 8.5 Hz), 7.57-7.61 (m, 1 H), 7.39 (s, 1 H), 7.34 (bs, 1 H), 6.93-7.01 (m, 4 H), 6.61-6.64 (m, 2 H), 4.62-4.80 (m, 1 H), 4.25-4.50 (m, 5 H), 3.15-3.20 (m, 1 H), 2.95-3.01 (m, 3 H), 2.75-2.87 (m, 4 H), 2.58-2.66 (m, 1 H), 2.36-2.49 (m, 2 H), 2.00-2.02 (m, 5 H), 1.69-1.88 (m, 3 H), 1.35-1.37 (m, 2  
15 H), 1.25-1.28 (m, 2 H); MS  $m/z$  ( $C_{37}H_{53}N_{11}O_{11}S+H$ )<sup>+</sup> 860.7; Anal. calcd for  $C_{37}H_{53}N_{11}O_{11}S$ : N, 17.92. Found: N, 14.34 (peptide content: 80%).

**5.47 Example 47: Acetyl-Pro-His-Ser-Cys-Asn-Ahp-Tyr-NH<sub>2</sub>**

This compound was prepared according to the procedures of Examples 1 and 2. The title compound (49.5 mg, 37 %) was isolated as a fine white powder. The NMR data indicated a mixture of two species in a ratio of about 80:20: <sup>1</sup>H NMR (300 MHz, DMSO-*d*<sub>6</sub>) δ 9.12 (bs, 1 H), 8.97-8.98 (m, 1 H), 8.26-8.29 (m, 1 H), 8.42 and 8.21 (d, 1 H, *J* = 7.6 Hz), 8.13 and 7.95 (d, 1 H, *J* = 7.8 Hz), 7.81 (d, 1 H, *J* = 8.5 Hz), 7.57-7.60 (m, 1 H), 7.39 (s, 1 H), 7.33 (bs, 2 H), 6.99-7.01 (app d, 2 H), 6.96 (s, 1 H), 6.93  
25 (s, 1 H), 6.61-6.64 (app d, 2 H), 4.60-4.79 (m, 1 H), 4.26-4.53 (m, 5 H), 3.11-3.21 (m, 1 H), 2.94-3.04 (m, 3 H), 2.73-2.88 (m, 3 H), 2.58-2.65 (m, 1 H), 2.39-2.49 (m, 2 H), 2.00-2.04 (m, 5 H), 1.70-1.91 (m, 3 H), 1.28-1.40 (m, 4 H), 1.05-1.20 (m, 4 H); MS  $m/z$  ( $C_{39}H_{57}N_{11}O_{11}S+H$ )<sup>+</sup> 888.8; Anal. calcd for  $C_{39}H_{57}N_{11}O_{11}S$ : N, 17.35. Found: N, 13.84 (peptide content: 80%).

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**5.48 Example 48: 3-(4-Hydroxyphenyl)propionyl-bAla-Pro-His-Ser-Cys-Asn-NH<sub>2</sub>**

This compound was prepared according to the procedures of Examples 1 and 2 with the following modifications: capping of the N-terminus was carried out using 3-(4-



hydroxyphenyl)propionic acid (3 eq), HBTU (3 eq), HOBT (3 eq) and NMM (6 eq). The title compound was isolated (61.1 mg, 53%) as a fine white powder. The NMR data indicated a mixture of two species in a ratio of about 75:25: <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>) δ 8.97 (bs, 1 H), 8.22-8.29 (m, 2 H), 8.10-8.13 (m, 1 H), 7.97 and 8.41 (d, 1 H, *J* = 7.2 Hz), 7.80 (app t, 1 H), 7.34-7.39 (m, 2 H), 7.08 (d, 2 H, *J* = 9.8 Hz), 6.92-6.98 (m, 3 H), 6.63-6.66 (app d, 2 H), 4.60-4.77 (m, 1 H), 4.27-4.50 (m, 5 H), 3.65-3.71 (m, 3 H), 3.39-3.49 (m, 3 H), 3.11-3.29 (m, 4 H), 2.94-3.04 (m, 1 H), 2.76-2.83 (m, 2 H), 2.37-2.46 (m, 3 H), 2.26-2.31 (m, 2 H), 1.97-2.08 (m, 1 H), 1.72-1.87 (m, 3 H); MS *m/z* (C<sub>33</sub>H<sub>46</sub>N<sub>10</sub>O<sub>10</sub>S+H)<sup>+</sup> 775.7; Anal. calcd for C<sub>33</sub>H<sub>46</sub>N<sub>10</sub>O<sub>10</sub>S: N, 18.08. Found: N, 12.76 (peptide content: 71%).

**5.49 Example 49: 8-(4-Fluorobenzylamino)suberoyl-β-Ala-Pro-His-Ser-Cys-Asn-NH<sub>2</sub>**

This compound was prepared according to the procedures of Examples 1 and 2 with the following modification: capping of the N-terminus was carried out using 8-(4-fluorobenzylamino)suberic acid (3 eq), HBTU (3 eq), HOBT (3 eq) and NMM (6 eq). The title compound was isolated as a fine white powder (70.8 mg, 53 %). The NMR data indicated a mixture of two species in a ratio of about 70:30: <sup>1</sup>H NMR (300 MHz, DMSO-d<sub>6</sub>) δ 8.97 (s, 1 H), 8.22-8.28 (m, 3 H), 8.11-8.14 (m, 1 H), 7.96-7.99 and 8.40-8.43 (m, 1 H), 7.74-7.80 (m, 1 H), 7.34-7.39 (m, 2 H), 7.24-7.29 (m, 2 H), 7.08-7.16 (m, 4 H), 6.92 (bs, 1 H), 4.29-4.50 (m, 7 H), 4.22 (d, 2 H, *J* = 5.9 Hz), 3.63-3.69 (m, 2 H), 3.44-3.52 (m, 2 H), 3.16-3.26 (m, 3 H), 2.97-3.02 (m, 1 H), 2.76-2.83 (m, 2 H), 2.38-2.49 (m, 3 H), 1.99-2.13 (m, 5 H), 1.72-1.89 (m, 3 H), 1.42-1.54 (m, 4 H), 1.20-1.24 (m, 4 H); MS *m/z* (C<sub>34</sub>H<sub>47</sub>N<sub>11</sub>O<sub>11</sub>S+H)<sup>+</sup> 818.6; Anal. Calcd for C<sub>39</sub>H<sub>56</sub>FN<sub>11</sub>O<sub>10</sub>S: N, 17.31. Found: N, 11.72 (peptide content: 68%).

**5.50 Example 50: m-dPEG-β-Ala-His-Ser-Cys-Asn-NH<sub>2</sub>**

This compound was prepared according to the procedures of Examples 1 and 2 with the following modification: capping of the N-terminus was carried out using NHS-m-dPEG<sup>TM</sup> (Quanta Biodesign, 1.9 eq), and TEA (9 eq). The title compound was isolated as a fine white powder (38 mg, 21%): <sup>1</sup>H NMR (300 MHz, D<sub>2</sub>O) δ 8.58 (d, 1 H, *J* = 1.2 Hz), 7.28 (d, 1 H, *J* = 0.8 Hz), 4.75-4.61 (m, 2 H), 4.53 (t, 1 H, *J* = 6.2 Hz), 4.44 (t, 1 H, *J* = 5.6 Hz), 4.34-4.29 (dd, 1 H, *J* = 5.1 Hz, 3.6 Hz), 3.82 (t, 2 H, *J* =

5.4 Hz), 3.71 (t, 2 H,  $J = 6.1$  Hz), 3.64-3.61 (m, 44 H), 3.58-3.54 (m, 4 H), 3.40 (t, 2 H,  $J = 6.7$  Hz), 3.63-3.69 (m, 2 H), 3.44-3.52 (m, 2 H), 3.16-3.26 (m, 3 H), 2.97-3.02 (m, 1 H), 3.32 (s, 3 H), 3.25 (d, 1 H,  $J = 5.9$  Hz), 3.18 (d, 1 H,  $J = 8.4$  Hz), 2.90 (d, 2 H,  $J = 6.2$  Hz), 2.83-2.56 (m, 3 H), 2.46 (t, 2 H,  $J = 6.1$  Hz), 2.25-2.15 (m, 1H), 1.92-  
5 1.87 (m, 1H), 1.83-1.76 (m, 1H); MS  $m/z$  ( $C_{50}H_{88}N_{10}O_{21}S + H$ )<sup>+</sup> 1198.1.

### 5.51 Example 51: DTPA conjugation to PHSCN

A solution of resin (Rink Amide AM resin) bound PHSCN (143mg, 0.36mmol/g, 0.051 mmol), p-SCN-benzyl DTPA (66mg, 0.103mmol), NNM (120μL, 10 2.1mmol), in anhydrous DMF (2mL) was agitated gently with nitrogen for 7 hours at room temperature, and the reaction mixture was filtered and washed with DMF once. This step was repeated twice more. The resin was washed three times with DMF, three times with methanol and three times with dichloromethane. The resulting resin bound compound was treated with TFA/TIS/water (95:2.5:2.5, 1 mL per 100 mg of  
15 resin) and agitated with nitrogen for 1 hour. The reaction mixture was filtered, the resin was washed once with TFA/TIS/water and three times with dichloromethane. The solvent was removed *in vacuo* and the resulting residue was triturated three times with ether. The resulting white powder was subjected to a prep HPLC purification according to the procedure of Example 2 to give 15.5 mg (0.014mmol, 27.7%) of  
20 desired product: <sup>1</sup>H NMR (300 MHz, D<sub>2</sub>O) δ 8.66 (s, 1H), 7.40 – 7.31 (m, 5H), 4.97-4.92 (dd,  $J = 4.92, 5.31$  Hz, 1H), 4.87-4.83 (m, 2H), 4.77-4.72 (m, 1H), 4.65-4.56 (m, 2H), 4.00-3.97 (m, 4H), 3.91-3.72 (m, 10H), 3.49-3.20 (m, 6H), 3.03-2.97 (m, 2H), 2.92-2.73 (m, 4H), 2.38 (m, 1H), 2.20 (m, 2H), 2.04-1.85 (m, 2H); MS  $m/z$  ( $C_{43}H_{61}N_{13}O_{17}S_2 + H$ )<sup>+</sup> 1096.8.

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### 5.52 Example 52: In<sup>3+</sup> chelation to DTPA-PHSCN

DTPA-PHSCN (20mg, 0.018mmol) was dissolved in 1mL of 0.1M AcOH(aq) solution and InCl<sub>3</sub> (40mg, 0.18mmol) was dissolved in 2mL of 0.02M HCl solution. The two solutions were combined and incubated for an hour at room  
30 temperature. The solvent was evaporated under reduced pressure and the resulting white powder was subjected to a prep HPLC purification according to the procedure of Example 2 to give 7.1 mg (0.00588mmol, 32.6%) of desired product: <sup>1</sup>H NMR (300 MHz, D<sub>2</sub>O) δ 8.66 (s, 1H), 7.41 – 7.31 (m, 5H), 4.97-4.92 (dd,  $J = 4.86, 5.46$  Hz,

1H), 4.86-4.84 (m, 2H), 4.76-4.72 (dd,  $J = 5.43, 2.88$  Hz, 1H), 4.61-4.57 (m, 2H), 4.04 (d,  $J = 17.22$  Hz, 1H), 3.90 (d,  $J = 5.82$  Hz, 2H), 3.81-3.60 (m, 5H), 3.50-3.19 (m, 10H), 3.14-2.73 (m, 8H), 2.39 (m, 1H), 2.16 (m, 1H), 2.04-1.91 (m, 2H); MS  $m/z$  ( $C_{43}H_{58}InN_{13}O_{17}S_2 + H$ )<sup>+</sup> 1208.8; Anal. calcd for  $C_{43}H_{58}InN_{13}O_{17}S_2$ : N, 15.07. Found: N, 11.58 (peptide content: 76.8%).

#### **5.53 Example 53: Acetyl-Pro-His-Ser-Cys-Asn-Gly-Gly-Gly-Lys-NH<sub>2</sub>**

This compound was prepared according to the procedures of Examples 1 and 2. The title compound was isolated (101 mg, 70%) as a fine white powder: <sup>1</sup>H NMR (300 MHz, D<sub>2</sub>O)  $\delta$  8.57 (d,  $J = 1.32$  Hz, 1H), 7.26 (s, 1H), 4.73-4.72 (m, 2H), 4.52 (t,  $J = 6.24$ , 1H), 4.43 (t,  $J = 5.25$  Hz, 1H), 4.27 (m, 2H), 3.94-3.91 (m, 8H), 3.81 (t,  $J = 5.40$  Hz, 2H), 3.57 (t,  $J = 6.78$  Hz, 1H), 3.31-3.24 (dd,  $J = 15.57, 5.67$  Hz, 1H), 3.18-3.09 (dd,  $J = 15.6, 8.64$  Hz, 2H), 2.96-2.83 (m, 4H), 2.80-2.73 (m, 2H), 2.20-2.16 (m, 1H), 2.05 (s, 3H), 1.94-1.57 (m, 6H), 1.42-1.34 (m, 2H); MS  $m/z$  ( $C_{37}H_{59}N_{15}O_{13}S + H$ )<sup>+</sup> 954.9.

#### **5.54 Example 54: Acetyl-Pro-His-Ser-Cys-Asn-Gly-Gly-Lys(Protoporphyrin)-NH<sub>2</sub>**

This compound was prepared according to procedures of Examples 2 and 3 with the following modifications: the 2% hydrazine treatment for the deprotection of ivDde on lysine was repeated 10 times and the amide formation on lysine was carried out with protoporphyrin IX (2 eq), PyBOP (2 eq), and NMM (6 eq). The title compound was isolated (7.6 mg, 5.49  $\mu$ M, 8.6%) as a fine dark red powder.; MS  $m/z$  ( $C_{67}H_{85}N_{17}O_{14}S + H$ )<sup>+</sup> 1385.6.

#### **5.55 Example 55: Ac-Tyr- $\beta$ -Ala-Pro-His-Ser-Cys-Asn-NH<sub>2</sub>**

This compound was prepared from Rink amide AM resin according to the procedures of Examples 1 and 2 and was isolated as a white, fluffy solid; ES MS  $m/z$  (M+H)<sup>+</sup> 832, (M+Na)<sup>+</sup> 854.

#### **5.56 Example 56: Ac-Pro-His-Ser-Cys-Asn- $\beta$ -Ala-Tyr-NH<sub>2</sub>**

This compound was prepared from Rink amide AM resin according to the procedures of Examples 1 and 2 and was isolated as a white, fluffy solid; ES MS  $m/z$   $(M+H)^+$  832,  $(M+Na)^+$  854.

5                    **5.57 Example 57: Ac-Pro-His-Ser-Cys-Asn-Gly-Gly-Lys-NH<sub>2</sub>**

This compound was prepared from Rink amide AM resin according to the procedures of Examples 1 and 2 and was isolated as a white, fluffy solid; ES MS  $m/z$   $(M+H)^+$  840

10                   **5.58 Example 58: Ac-Pro-His-Ser-Cys-Asn-Gly-Gly-Lys(biotin)-NH<sub>2</sub>**

This compound was prepared from Rink amide AM resin according to the procedures of Examples 1 and 2 and was isolated as a white, fluffy solid; ES MS  $m/z$   $(M+H)^+$  1066.5.

15                   Finally, it should be noted that there are alternative ways of implementing the present invention. Accordingly, the present embodiments are to be considered as illustrative and not restrictive, and the invention is not to be limited to the details given herein, but may be modified within the scope and equivalents of the appended claims. All publications and patents cited herein are incorporated by reference in their  
20                   entirety.